

Protein Sequence Searches - February 2005

All of the sequence databases on ABSS have recently been updated.

- Please note that the curators of the UniProt database have purged some temporary accession numbers from the most recent version of UniProt. These sequences have been assigned new permanent accession numbers. The new UniProt record may not contain the previous temporary accession number.
- If you encounter an accession number from an older search run against UniProt (results file extension **.rup**) that can no longer be found in the database, the permanent record with the new accession number can be found by searching the old accession number in the UniProt Protein Archive database (UniPARC) at:

<http://www.pir.uniprot.org/database/archive.shtml>

If you have any questions regarding this information or your results, please contact any STIC searcher.

When submitting sequence search results for scanning into IFW, please include a copy of this attachment to assist any future Examiners or members of the public who may encounter UniProt temporary accession numbers.

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160352

From: Swope, Sheridan
Sent: Monday, July 25, 2005 2:40 PM
To: STIC-Biotech/ChemLib
Subject: 10/726,967

For 10/726,967, pls search:

(A)--Claim 1(18)

SID 3: oligo search against the NT & AA data bases.

(B)

SID 52 against the NT & AA data bases.

For any hits that are 100% identical, pls align with:
residues 22-41 of SID 1,

CRF

If you have questions, pls ask me or David Schreiber.

Thanks

Sheridan Swope, Ph.D.
Patent Examiner, AU 1656
Recombinant Enzymes
571-272-0943 (voice)
E02B71 Remsen Bld (Office)
E03C70 Remsen Bld (Mailbox)

STAFF USE ONLY

Searcher: _____
Searcher Phone: 2-_____
Date Searcher Picked up: _____
Date Completed: _____
Searcher Prep/Rev. Time: _____
Online Time: _____

Type of Search

NA#: _____ AA#: _____
Interference: _____ SPDI: _____
S/L: _____ Oligomer: _____
Encode/Transl: _____
Structure#: _____ Text: _____
Inventor: _____ Litigation: _____

Vendors and cost where applicable

STN: _____
DIALOG: _____
QUESTEL/ORBIT: _____
LEXIS/NEXIS: _____
SEQUENCE SYSTEM: _____
WWW/Internet: _____
Other(Specify): _____

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(uspto)

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: July 26, 2005, 16:28:49 ; Search time 43 Seconds
(without alignments)
27.776 Million cell updates/sec

Title: US-10-726-967A-3

Perfect score: 16

Sequence: 1 TQHGRLPLRSGLGCA 16

Scoring table: OLIGO

Gapop 60.0 , Gapext 60.0

Searched: 513545 seqs, 74649064 residues

Word size : 0

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database : Issued Patents AA:*

1: /cgn2_6/prodata/1/iaa/5A_COMB.pep:*\n2: /cgn2_6/prodata/1/iaa/5B_COMB.pep:*\n3: /cgn2_6/prodata/1/iaa/6A_COMB.pep:*\n4: /cgn2_6/prodata/1/iaa/6B_COMB.pep:*\n5: /cgn2_6/prodata/1/iaa/6C_COMB.pep:*\n6: /cgn2_6/prodata/1/iaa/6D_COMB.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	24	US-09-724-566A-47	Sequence 47, Appl
2	16	100.0	419	US-09-724-566A-57	Sequence 57, Appl
3	16	100.0	419	US-09-471-669A-57	Sequence 57, Appl
4	16	100.0	420	US-09-724-566A-60	Sequence 60, Appl
5	16	100.0	420	US-09-471-669A-60	Sequence 60, Appl
6	16	100.0	428	US-09-548-372D-51	Sequence 51, Appl
7	16	100.0	428	US-09-548-367D-51	Sequence 51, Appl
8	16	100.0	428	US-09-551-853D-51	Sequence 51, Appl
9	16	100.0	428	US-09-548-367D-51	Sequence 51, Appl
10	16	100.0	428	US-09-548-367D-51	Sequence 51, Appl
11	16	100.0	428	US-09-794-927A-51	Sequence 51, Appl
12	16	100.0	428	US-09-548-373D-51	Sequence 51, Appl
13	16	100.0	428	US-09-795-847B-51	Sequence 51, Appl
14	16	100.0	428	US-09-869-414-51	Sequence 51, Appl
15	16	100.0	428	US-09-548-366F-51	Sequence 51, Appl
16	16	100.0	428	US-09-548-368D-51	Sequence 51, Appl
17	16	100.0	428	US-09-794-925A-51	Sequence 51, Appl
18	16	100.0	428	US-09-724-566A-74	Sequence 74, Appl
19	16	100.0	431	US-09-471-669A-74	Sequence 74, Appl
20	16	100.0	433	US-09-548-372D-26	Sequence 26, Appl
21	16	100.0	433	US-09-548-367D-26	Sequence 26, Appl
22	16	100.0	433	US-09-551-853D-26	Sequence 26, Appl
23	16	100.0	433	US-09-416-801B-26	Sequence 26, Appl
24	16	100.0	433	US-09-548-376D-26	Sequence 26, Appl
25	16	100.0	433	US-09-794-927A-26	Sequence 26, Appl
26	16	100.0	433	US-09-548-373D-26	Sequence 26, Appl
27	16	100.0	433	US-09-795-847B-26	Sequence 26, Appl

28	16	100.0	433	US-09-869-414-26	Sequence 26, Appl
29	16	100.0	433	US-09-548-366F-26	Sequence 26, Appl
30	16	100.0	433	US-09-548-368D-26	Sequence 26, Appl
31	16	100.0	433	US-09-794-925A-26	Sequence 26, Appl
32	16	100.0	433	US-09-806-194A-26	Sequence 26, Appl
33	16	100.0	433	US-09-548-372D-53	Sequence 53, Appl
34	16	100.0	434	US-09-548-367D-53	Sequence 53, Appl
35	16	100.0	434	US-09-551-853D-53	Sequence 53, Appl
36	16	100.0	434	US-09-416-901B-53	Sequence 53, Appl
37	16	100.0	434	US-09-548-376D-53	Sequence 53, Appl
38	16	100.0	434	US-09-794-927A-53	Sequence 53, Appl
39	16	100.0	434	US-09-548-372D-53	Sequence 53, Appl
40	16	100.0	434	US-09-795-847B-53	Sequence 53, Appl
41	16	100.0	434	US-09-869-414-53	Sequence 53, Appl
42	16	100.0	434	US-09-548-366F-53	Sequence 53, Appl
43	16	100.0	434	US-09-548-368D-53	Sequence 53, Appl
44	16	100.0	434	US-09-794-925A-53	Sequence 53, Appl
45	16	100.0	446	US-09-548-372D-22	Sequence 22, Appl

ALIGNMENTS

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RESULT 1
US-09-724-566A-47
; Sequence 47, Application US/09724566A
; Patent No. 6627739
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Bassi, Guribai
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6627739mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Simha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
; FILE REFERENCE: 228-US-NEWC2
; CURRENT FILING DATE: 2000-11-28
; PRIOR FILING DATE: 1999-06-15
; PRIOR FILING DATE: 2000-02-10
; PRIOR APPLICATION NUMBER: 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 47
; LENGTH: 24
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-724-566A-47

Query Match      100.0%; Score 16; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 86-10;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy      1 TQHGRLPLRSGLGCA 16
Db      1 TQHGRLPLRSGLGCA 16

RESULT 2
US-09-724-566A-57
; Sequence 57, Application US/09724566A
; Patent No. 6627739
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
```

```

/ APPLICANT: Bael, Gurigbal
/ APPLICANT: Doane, Minh Tam
/ APPLICANT: Frigon, No. 6627739mand
/ APPLICANT: John, Varghese
/ APPLICANT: Power, Michael
/ APPLICANT: Sinha, Sukanto
/ APPLICANT: Tatsuno, Gwen
/ APPLICANT: Tung, Jay
/ APPLICANT: Wang, Shuwen
/ APPLICANT: McConlogue, Lisa
/ TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
/ FILE REFERENCE: 228-US-NEWC2
/ CURRENT APPLICATION NUMBER: US/09/724,566A
/ PRIOR FILING DATE: 2000-11-28
/ PRIOR APPLICATION NUMBER: US 09/501,708
/ PRIOR FILING DATE: 2000-02-10
/ PRIOR APPLICATION NUMBER: 60/119,571
/ PRIOR FILING DATE: 1999-02-10
/ PRIOR APPLICATION NUMBER: 60/139,172
/ PRIOR FILING DATE: 1999-06-15
/ NUMBER OF SEQ ID NOS: 104
/ SOFTWARE: FaSTSeq for Windows Version 4.0
/ SEQ ID NO 57
/ LENGTH: 419
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ US-09-724-566A-57
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Query Match      100.0%; Score 16; DB 4; Length 419;
Best Local Similarity 100.0%; Pred. No. 1,1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1  TQHGIRLPRLSGGGA 16
Db      22  TQHGIRLPRLSGGGA 37
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RESULT 3
/ US-09-471-669A-57
/ Sequence 57, Application US/09471669A
/ Patent No. 6830918
/ GENERAL INFORMATION:
/ APPLICANT: Anderson, John P.
/ APPLICANT: Bael, Gurigbal
/ APPLICANT: Doane, Minh Tam
/ APPLICANT: Frigon, No. 6830918mand
/ APPLICANT: John, Varghese
/ APPLICANT: Power, Michael
/ APPLICANT: Sinha, Sukanto
/ APPLICANT: Tatsuno, Gwen
/ APPLICANT: Tung, Jay
/ APPLICANT: Wang, Shuwen
/ APPLICANT: McConlogue, Lisa
/ TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
/ FILE REFERENCE: 015270-006430US
/ CURRENT APPLICATION NUMBER: US/09/471,669A
/ PRIOR FILING DATE: 1999-12-24
/ PRIOR APPLICATION NUMBER: US 60/114,408
/ PRIOR FILING DATE: 1998-12-31
/ PRIOR APPLICATION NUMBER: US 60/119,571
/ PRIOR FILING DATE: 1999-02-10
/ PRIOR APPLICATION NUMBER: US 60/139,172
/ PRIOR FILING DATE: 1999-06-15
/ NUMBER OF SEQ ID NOS: 108
/ SOFTWARE: PatentIn Ver. 2.1
/ SEQ ID NO 57
/ LENGTH: 419
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ US-09-471-669A-57
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Query Match      100.0%; Score 16; DB 4; Length 419;
Best Local Similarity 100.0%; Pred. No. 1,1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      1  TQHGIRLPRLSGGGA 16
Db      22  TQHGIRLPRLSGGGA 37
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RESULT 4
/ US-09-724-566A-60
/ Sequence 60, Application US/09724566A
/ Patent No. 6627739
/ GENERAL INFORMATION:
/ APPLICANT: Anderson, John P.
/ APPLICANT: Bael, Gurigbal
/ APPLICANT: Doane, Minh Tam
/ APPLICANT: Frigon, No. 6627739mand
/ APPLICANT: John, Varghese
/ APPLICANT: Power, Michael
/ APPLICANT: Sinha, Sukanto
/ APPLICANT: Tatsuno, Gwen
/ APPLICANT: Tung, Jay
/ APPLICANT: Wang, Shuwen
/ APPLICANT: McConlogue, Lisa
/ TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
/ FILE REFERENCE: 228-US-NEWC2
/ CURRENT APPLICATION NUMBER: US/09/724,566A
/ PRIOR FILING DATE: 2000-11-28
/ PRIOR APPLICATION NUMBER: US 09/501,708
/ PRIOR FILING DATE: 2000-02-10
/ PRIOR APPLICATION NUMBER: 60/119,571
/ PRIOR FILING DATE: 1999-02-10
/ PRIOR APPLICATION NUMBER: 60/139,172
/ PRIOR FILING DATE: 1999-06-15
/ NUMBER OF SEQ ID NOS: 104
/ SOFTWARE: FaSTSeq for Windows Version 4.0
/ SEQ ID NO 60
/ LENGTH: 420
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ US-09-724-566A-60
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Query Match      100.0%; Score 16; DB 4; Length 420;
Best Local Similarity 100.0%; Pred. No. 1,1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1  TQHGIRLPRLSGGGA 16
Db      22  TQHGIRLPRLSGGGA 37
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```

RESULT 5
/ US-09-471-669A-60
/ Sequence 60, Application US/09471669A
/ Patent No. 6830918
/ GENERAL INFORMATION:
/ APPLICANT: Anderson, John P.
/ APPLICANT: Bael, Gurigbal
/ APPLICANT: Doane, Minh Tam
/ APPLICANT: Frigon, No. 6830918mand
/ APPLICANT: John, Varghese
/ APPLICANT: Power, Michael
/ APPLICANT: Sinha, Sukanto
/ APPLICANT: Tatsuno, Gwen
/ APPLICANT: Tung, Jay
/ APPLICANT: Wang, Shuwen
/ APPLICANT: McConlogue, Lisa
/ TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
/ FILE REFERENCE: 015270-006430US
/ CURRENT APPLICATION NUMBER: US/09/471,669A
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; CURRENT FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: Patentn Ver. 2.1
; SEQ ID NO 60
; LENGTH: 420
; TYPE: PRT
; ORGANISM: Homo sapiens
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-471-669A-60

Query Match          100.0%; Score 16; DB 4; Length 420;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGCA 16
   |||||
Db 22 TOHGIRLPLRSGGCA 37

RESULT 6
US-09-548-372D-51
; Sequence 51, Application US/09548372D
; Patent No. 6420534
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/62801
; CURRENT FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentn version 3.1
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-372D-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGCA 16
   |||||
Db 22 TOHGIRLPLRSGGCA 37

RESULT 7
US-09-548-367D-51
; Sequence 51, Application US/09548367D
; Patent No. 6440698
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280H
; CURRENT FILING DATE: US/09/548,367D
; CURRENT FILING DATE: 2000-04-12
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; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentn version 3.1
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-367D-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGCA 16
   |||||
Db 22 TOHGIRLPLRSGGCA 37

RESULT 8
US-09-551-853D-51
; Sequence 51, Application US/09551853D
; Patent No. 6500667
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280L
; CURRENT FILING DATE: 2000-04-18
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentn version 3.1
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-551-853D-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGCA 16
   |||||
Db 22 TOHGIRLPLRSGGCA 37

RESULT 9
US-09-416-901B-51
; Sequence 51, Application US/09416901B
; Patent No. 669671
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: THEREOF
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FILE REFERENCE: 29915/6280A
CURRENT APPLICATION NUMBER: US/09/416,901B
CURRENT FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 72
SOFTWARE: Patentin version 3.1
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-416-901B-51
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Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1 TOHGIRLPLRSGGGA 16
        |||||||
Db      22 TOHGIRLPLRSGGGA 37
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```
RESULT 10
US-09-548-376D-51
Sequence 51, Application US/09548376D
Patent No. 6706485
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR
TITLE OF INVENTION: AND USES
FILE REFERENCE: 29915/6280F
CURRENT APPLICATION NUMBER: US/09/548,376D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: Patentin version 3.1
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-376D-51
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Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1 TOHGIRLPLRSGGGA 16
        |||||||
Db      22 TOHGIRLPLRSGGGA 37
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```
RESULT 11
US-09-794-927A-51
Sequence 51, Application US/09794927A
Patent No. 6727074
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```
GENERAL INFORMATION:
APPLICANT: Gurney et al.
TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
TITLE OF INVENTION: THEREFOR
FILE REFERENCE: 29915/6280FG
CURRENT APPLICATION NUMBER: US/09/794,927A
CURRENT FILING DATE: 2001-02-27
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 74
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-794-927A-51
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Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1 TOHGIRLPLRSGGGA 16
        |||||||
Db      22 TOHGIRLPLRSGGGA 37
```

```
RESULT 12
US-09-548-373D-51
Sequence 51, Application US/09548373D
Patent No. 6737510
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
TITLE OF INVENTION: THEREOF
FILE REFERENCE: 29915/6280B
CURRENT APPLICATION NUMBER: US/09/548,373D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: Patentin version 3.1
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-373D-51
```

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Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1 TOHGIRLPLRSGGGA 16
        |||||||
Db      22 TOHGIRLPLRSGGGA 37
```

```
RESULT 13
US-09-795-847B-51
; Sequence 51, Application US/09795847B
; Patent No. 6753163
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280DE
; CURRENT APPLICATION NUMBER: US/09/795,847B
; CURRENT FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-795-847B-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGIGGA 16
        |||||||
Db      22 TOHGIRLPLRSGIGGA 37

RESULT 14
US-09-869-414-51
; Sequence 51, Application US/09869414
; Patent No. 6790610
; GENERAL INFORMATION:
; APPLICANT: Bienkowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M
; CURRENT APPLICATION NUMBER: US/09/869,414
; CURRENT FILING DATE: 2001-06-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-869-414-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGIGGA 16
        |||||||
Db      22 TOHGIRLPLRSGIGGA 37
```

```
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-869-414-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGIGGA 16
        |||||||
Db      22 TOHGIRLPLRSGIGGA 37

RESULT 15
US-09-548-366F-51
; Sequence 51, Application US/09548366F
; Patent No. 6797487
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280J
; CURRENT APPLICATION NUMBER: US/09/548,366F
; CURRENT FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-366F-51

Query Match          100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGIGGA 16
        |||||||
Db      22 TOHGIRLPLRSGIGGA 37

Search completed: July 26, 2005, 16:38:50
Job time : 44 secs
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CC therapy. (1) can be used for producing preparations of homogeneously
CC processed BACE that may be used for e.g. studying or treating diseases
CC such as Alzheimer's disease or Down's syndrome. The human BACE1 gene is
CC located on chromosome 11, more specifically to 11q23.2-23.3. The present
CC sequence represents the human BACE1 isoform A 22-37 prodomain amino acid
CC sequence, which is used in the exemplification of the present invention.
SQ Sequence 16 AA;

Query Match 100.0%; Score 16; DB 8; Length 16;
Best Local Similarity 100.0%; Pred. No. 7e-09;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DQ 1 TOHGIRLPRLRSGLGGA 16
1 TOHGIRLPRLRSGLGGA 16

RESULT 2
ADC81580
ID ADC81580 standard; protein; 425 AA.
AC ADC81580;
XX
XX 01-JAN-2004 (first entry)
DT
XX
XX Beta-secretase zymogen (pbsz) amino acid sequence SEQ ID NO:3.
DE
XX human; BACE; modification; Pro33lys; pro-enzyme.
KM
XX Unidentified.
OS
XX WO2003072733-A2.
PN
XX 04-SEP-2003.
PD
XX 21-FEB-2003; 2003WO-US005508.
PF
XX 21-FEB-2002; 2002US-0356651P.
PR
XX (PMAA) PHARMACIA & UPJOHN CO.
PA
XX Chou K, Howe JW;
PI
XX WPI; 2003-712719/67.
DR
XX
XX BACE polypeptides having Pro33lys modification, useful in determining
PT possible mutations, which will inhibit enzyme activity, and in
PT determining potential active site for target molecules.
PS Disclosure; Fig 3; 38pp; English.

CC The present invention describes an isolated polypeptide (1) comprising or
CC consisting of a fully defined sequence of 432 amino acids (see ADC81561),
CC and comprising human BACE having the modification Pro33lys. Also
CC described: (1) a composition comprising an active human BACE enzyme
CC comprising the pro-enzyme sequence of BACE having the modification
CC Pro33lys; (2) an isolated polynucleotide comprising a sequence encoding
CC (1); (3) an isolated polynucleotide comprising a sequence encoding
CC nucleotides 70-1365 of a 1355-bp sequence (see ADC81562); (4) an
CC expression vector comprising the polynucleotide of (2); or a
CC polynucleotide sequence encoding a Pro33lys-BACE polypeptide, where the
CC expression vector can produce the Pro33lys-BACE polypeptide, where the
CC in a compatible host cell, when cultured under conditions that allow
CC production; (5) a recombinant host cell comprising the expression vector;
CC and (6) producing a (active) Pro33lys-BACE polypeptide, the BACE
CC polypeptide having Pro33lys modification may be used in determining
CC possible mutations, which will inhibit enzyme activity, and in
CC determining potential active site for target molecules. The vector
CC comprising the BACE polynucleotide is useful for producing recombinant
CC BACE polypeptides having Pro33lys modification. The present sequence
CC represents a beta-secretase zymogen amino acid sequence, which is used in
CC the exemplification of the present invention.

XX
SQ Sequence 425 AA;

Query Match 100.0%; Score 16; DB 7; Length 425;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DQ 1 TOHGIRLPRLRSGLGGA 16
1 TOHGIRLPRLRSGLGGA 16

RESULT 3
AAU07219
ID AAU07219 standard; protein; 428 AA.
AC AAU07219;
XX
XX 24-OCT-2001 (first entry)
DT
XX
XX Human aspartyl protease 2b delatM (HuAsp-2bdeletM).DE
XX Human; aspartyl protease 1; Asp-1; neuroprotective;
KW aspartyl protease 2; Asp2; amyloid protein precursor; App;
KW beta-secretase; Alzheimer's disease; HuAsp-2bdeletM.
XX
XX Homo sapiens.OS

Key Location/Qualifiers
FH Misc-difference 1 /note= "Encoded by NNN"
FT Misc-difference 2 /note= "Encoded by NNC"
FT Misc-difference 61 /note= "Encoded by NNC"
FT Misc-difference 62 /note= "Encoded by NNC"
FT Misc-difference 121 /note= "Encoded by NNC"
FT Misc-difference 122 /note= "Encoded by NNC"
FT Misc-difference 181 /note= "Encoded by NNC"
FT Misc-difference 182 /note= "Encoded by NNC"
FT Misc-difference 241 /note= "Encoded by NNC"
FT Misc-difference 242 /note= "Encoded by NNC"
FT Misc-difference 301 /note= "Encoded by NNC"
FT Misc-difference 302 /note= "Encoded by NNC"
FT Misc-difference /note= "Encoded by NNT"

WO200149097-A2.
12-JUL-2001.
09-MAY-2001; 2001WO-IB000797.
PR 09-MAY-2001; 2001WO-IB000797.
XX
XX (BIEN/) BIENKOWSKI M J.
PA (GURN/) GURNEY M E.
PA (HEIN/) HEINRIKSON R L.
PA (PARO/) PARODI L A.
PA (YANR/) YAN R.
PI
XX
XX Blenkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
DR WPI; 2001-502548/55.
DR N-PSDB; AAS11732.
XX

PT Novel purified polypeptide comprising fragment of mammalian aspartyl
PT protease 2, lacking Asp2 transmembrane domain and retaining beta
PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
PT activity.
PS Claim 149; Page 167-168; 185pp; English.
XX
CC The invention relates to a novel purified polypeptide comprising a
CC fragment of mammalian aspartyl protease 2 (Asp2) protein which lacks the
CC Asp2 transmembrane domain and the Asp2 protein, and where the polypeptide
CC and the fragment retain the beta-secretase activity of the mammalian Asp2
CC protein. Also included is an isoform of amyloid protein precursor (APP)
CC comprising the amino acid sequence of a APP or its fragment containing an
CC APP cleavage site recognizable by a mammalian beta-secretase, and further
CC comprising two lysine residues at the carboxyl terminus of the amino acid
CC sequence of the mammalian APP or APP fragment. The polypeptides are used
CC for assaying for modulators of beta-secretase activity; identifying
CC agents that inhibit the APP processing activity of human Asp2 aspartyl
CC protease (Hu-Asp2); identifying agents that modulate the activity of Asp2
CC; and for reducing cellular production of amyloid beta (Abeta) from APP.
CC Agents identified by the above methods are useful for treating
CC Alzheimer's disease; and for identifying modulators of amyloid-beta
CC (Abeta) peptide production, for use in designing therapeutics for the
CC treatment or prevention of Alzheimer's disease. Probes and primers
CC derived from Asp nucleic acid sequences are useful for detecting Hu-Asp
CC nucleic acids in in vitro assays and in Northern and Southern blots. The
CC present sequence represents the amino acid sequence of human Asp-2b delta
CC TM construct which lacks the transmembrane domain. This construct was
CC used for bacterial expression and purification of human Asp2b
CC
XX
SQ Sequence 428 AA;
Query Match 100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TQHGIRLPLRSGIGGA 16
DB 22 TQHGIRLPLRSGIGGA 37
RESULT 4
AAE10646
ID AAE10646 standard; protein; 428 AA.
AC AAE10646;
XX
XX 10-DEC-2001 (first entry)
DT Human-Asp 2(b) protein lacking transmembrane domain.
DE Human: aspartyl protease 2b; Asp2b; amyloid precursor protein; APP;
XX Alzheimer's disease; AD; dementia; neurofibrillary tangle; gliosis;
KW amyloid plaque; neuronal loss; proteolytic; neurotropic; neuroprotective.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX GB2357767-A.
PN
XX
PD 04-JUL-2001.
XX
PF 22-SEP-2000; 2000GB-00023315.
XX
XX 23-SEP-1999; 99US-00404133.
PR 23-SEP-1999; 99US-0155493P.
PR 23-SEP-1999; 99WO-US020881.
PR 13-OCT-1999; 99US-00416901.
PR 06-DEC-1999; 99US-0169232P.
XX
XX (PHAA) PHARMACIA & UPJOHN CO.
PA
XX
PI Bienkowski MJ, Gurney M;

XX
DR WPI; 2001-444208/48.
DR N-PSDB; AAD17895.
XX
XX Polypeptide comprising fragments of human aspartyl protease with amyloid
PT precursor protein processing activity and alpha-secretase activity, for
PT identifying modulators useful in treating Alzheimer's disease.
XX
PS Example 10; Page 138-139; 187pp; English.
XX
CC The patent discloses human aspartyl protease 1 (hu-Asp1) or modified Asp1
CC proteins which lack transmembrane domain or amino terminal domain or
CC cytoplasmic domain and retains alpha-secretase activity and amyloid
CC protein precursor (APP) processing activity. The proteins of the
CC invention are useful for assaying hu-Asp1 alpha-secretase activity, which
CC in turn is useful for identifying modulators of hu-Asp1 alpha-secretase
CC activity, where modulators that increase hu-Asp1 alpha-secretase activity
CC are useful for treating Alzheimer's disease (AD) which causes progressive
CC dementia with consequent formation of amyloid plaques, neurofibrillary
CC tangles, gliosis and neuronal loss. Hu-Asp1 protease substrate is useful
CC for assaying hu-Asp1 proteolytic activity, by contacting hu-Asp1 protein
CC with the substrate under acidic conditions and determining the level of
CC hu-Asp1 proteolytic activity. The present sequence is human Asp 2(b)
CC protein lacking a transmembrane (TM) domain. This sequence is generated
CC by the deletion of the C-terminal TM domain and intracellular domains of
CC human Asp 2(b) protein
CC
XX
SQ Sequence 428 AA;
Query Match 100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TQHGIRLPLRSGIGGA 16
DB 22 TQHGIRLPLRSGIGGA 37
RESULT 5
AAE06891
ID AAE06891 standard; protein; 428 AA.
AC AAE06891;
XX
XX 23-OCT-2001 (first entry)
DT Human-Asp2(b) deltatM protein.
DE Human: aspartyl protease 2b; Asp 2b; beta-amyloid precursor protein; APP;
KW beta-secretase; Alzheimer's disease; dementia; amyloid plaque; gliosis;
KW neurofibrillary tangle; neuronal loss; amyloid-beta peptide; neurotropic;
KW neuroprotective; antisense therapy; Asp2(b) deltatM protein;
XX gene therapy.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX WO200150829-A2.
PN
XX
PD 19-JUL-2001.
XX
PF 09-MAY-2001; 2001WO-1B000799.
XX
XX 09-MAY-2001; 2001WO-1B000799.
PR
XX
XX (BIEN/) BIENKOWSKI M J.
PA (GURN/) GURNEY M B.
PA (HEIN/) HEINRIKSON R L.
PA (PARO/) PARODI L A.
PA (YANR/) YAN R.
XX
XX Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
PI

DR WPI: 2001-483072/52.
DR N-PSDB; AAD13276.

PT Novel purified polypeptide comprising fragment of mammalian aspartyl
PT protease 2, lacking Asp2 transmembrane domain and retaining beta
PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
PT activity.

PS Claim 149; Page 167-168; 185pp; English.

XX
XX The invention relates to human aspartyl proteases (Hu-Asp), beta-amyloid
CC precursor protein (APP) isoforms and their corresponding DNA molecules.
CC Human aspartyl proteases can act as beta-secretase proteases useful for
CC treating Alzheimer's disease. APP isoforms are useful for identifying
CC modulators of amyloid-beta peptide production, for use in designing
CC therapeutics for the treatment and prevention of Alzheimer's disease,
CC and neuronal loss. APP isoforms are also used in methods for identifying
CC inhibitors and modulators of human Asp2 activity. The invention relates
CC to a method for identifying agents that modulate the activity of human
CC aspartyl protease Asp2. Amyloid-beta peptides obtained from APP are used
CC as a means to screen in cellular assays for the inhibitors of beta- and
CC gamma-secretase. Hu-Asp DNA fragments are useful as probes or primers in
CC polymerase chain reactions (PCR). The probes are useful for detecting Hu-
CC Asp nucleic acids in in vitro assays and in Northern and Southern blots.
CC The present sequence is Human aspartyl protease 2b (Hu-Asp2b) deltaTM
CC protein which is obtained by the deletion of C-terminal transmembrane and
CC intracellular domains of Hu-Asp2b. Human Asp2b has beta-secretase
CC activity

XX Sequence 428 AA;

Query Match 100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TOHGIRLPRLRSGLGGA 16
|||
DB 22 TOHGIRLPRLRSGLGGA 37

RESULT 6
AAE02598
ID AAE02598 standard; protein; 428 AA.
XX
AC AAE02598;

DT 10-AUG-2001 (first entry)
XX

DE Human aspartyl protease 2 (b) delta TM protein.

XX Human; alpha-secretase; amyloid precursor protein; APP; therapy;
KW Alzheimer's disease; antialzheimer's; aspartyl protease 2; Asp 2;
KW beta-secretase; chromosome 11q23.3-24.1; mutant; mutein.

OS Homo sapiens.
OS Synthetic.

PN WO200123533-A2.

PD 05-APR-2001.

PF 22-SEP-2000; 2000WO-US026080.

PR 23-SEP-1999; 99US-0155493P.

PR 23-SEP-1999; 99WO-US020881.

PR 13-OCT-1999; 99US-00416901.

PR 06-DEC-1999; 99US-0169232P.

PA (PHAA) PHARMACIA & UPJOHN CO.

XX Gurney M, Bienkowski MJ;
XX
PI
XX

DR WPI: 2001-290516/30.
DR N-PSDB; AAD06768.

PT Enzymes that cleave the alpha-secretase site of the amyloid precursor
PT protein, useful for the treatment of Alzheimer's disease.

PS Example 10; Page 166-167; 189pp; English.

XX
XX The present invention relates to enzymes for cleaving the alpha-
CC secretase site of the amyloid precursor protein (APP) and methods of
CC identifying those enzymes. The methods may be used to identify enzymes
CC that may be used to cleave the alpha-secretase cleavage site of the APP
CC protein. The enzymes may be used to treat or modulate the progress of
CC Alzheimer's disease. The present sequence is human aspartyl protease 2
CC (Asp 2) (b) delta TM protein. The Asp 2 gene is located on chromosome
CC 11q23.3-24.1. The Asp 2 has beta-secretase protease activity

XX Sequence 428 AA;

Query Match 100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TOHGIRLPRLRSGLGGA 16
|||
DB 22 TOHGIRLPRLRSGLGGA 37

RESULT 7
AAU06620
ID AAU06620 standard; protein; 428 AA.
XX
AC AAU06620;

DT 24-OCT-2001 (first entry)
XX

DE Human-pro-Asp 2 (b) delta TM.

XX Human; Aspartyl protease; beta-secretase; neurotropic; ASP2;
KW neuroprotective; amyloid protein precursor; APP; Alzheimer's disease;
KW amyloid-beta; Abeta; Human-pro-Asp 2 (b) delta TM; mutant; mutein.

OS Homo sapiens.
OS Synthetic.

PN WO200149098-A2.

PD 12-JUL-2001.

PF 09-MAY-2001; 2001WO-IB000798.

PR 09-MAY-2001; 2001WO-IB000798.

PA (BIEN/) BIENKOWSKI M J.

PA (GURN/) GURNEY M E.

PA (HEIN/) HEINRIKSON R L.

PA (PARO/) PARODI L A.

PA (YANR/) YAN R.

PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;

DR WPI: 2001-502549/55.

XX
XX Novel purified polypeptide comprising fragment of mammalian aspartyl
PT protease 2, lacking Asp2 transmembrane domain and retaining beta
PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
PT activity.

PS Claim 149; Page 167-168; 185pp; English.

XX The invention relates to a purified polypeptide comprising a fragment of
CC mammalian aspartyl protease (Asp)2 protein which lacks the Asp2
CC transmembrane domain and the Asp2 protein, and where the polypeptide and

CC the fragment retain the beta-secretase activity of the mammalian Asp2
CC protein. The invention also details polynucleotides for the Asp proteins
CC and vectors expressing them, and a polypeptide (isoform of amyloid
CC protein precursor (APP)) comprising the amino acid sequence of an APP or
CC its fragment containing an APP cleavage site recognizable by a mammalian
CC beta-secretase, and further comprising two lysine residues at the
CC carboxyl terminus of the amino acid sequence of the mammalian APP or APP
CC fragment. Also included in the invention are methods of identifying
CC modulators or inhibitors of Asp2. Modulators and inhibitors of Asp2 are
CC useful for treating Alzheimer's disease. APP is useful in methods for
CC identifying inhibitors or modulators of human Asp2 activity and amyloid-
CC beta (Abeta) peptide production. APP is also useful in designing
CC therapeutics for the treatment or prevention of Alzheimer's disease. APP
CC comprising the APP-Sw-beta-secretase peptide sequence (NLDA), which is
CC associated with increased levels of Abeta processing is useful in assays
CC relating the Alzheimer's research. The expression vector is useful for
CC recombinantly expressing APP. Nucleic acids that hybridize to APP
CC oligonucleotides are useful as probes or primers. The probes are useful
CC for detecting Hu-Asp nucleic acids in *in vitro* assays and in Northern and
CC Southern blots. The present sequence is Human-pro-Asp 2(b) delta TM
CC protein, which lacks the C-terminal transmembrane domain
CC
XX
SQ Sequence 428 AA;

Query Match 100.0%; Score 16; DB 4; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TQHGIRLPRLSGGGA 16
|||
22 TQHGIRLPRLSGGGA 37

RESULT 8
AB878607
ID AB878607 standard; protein; 428 AA.

AC AB878607;

DT 16-JUN-2002 (first entry)

DE Human Asp-2(b) deltatm protein sequence SEQ ID NO:51.

XX Human; Asp-1; Asp-2; aspartyl protease; Alzheimer's disease; proteolytic;
XX Chromosome 11q23.3-24.1.

XX Homo sapiens.

XX GB2367060-A.

XX 27-MAR-2002.

XX 29-OCT-2001; 2001GB-00025934.

XX 23-SEP-1999; 99US-00404133.

XX 23-SEP-1999; 99US-0155493P.

XX 23-SEP-1999; 99WO-US020881.

XX 13-OCT-1999; 99US-00416901.

XX 06-DEC-1999; 99US-0169232P.

XX 22-SEP-2000; 2000GB-00023315.

XX (PHAA) PHARMACIA & UPJOHN CO.

XX Bienkowski MJ, Gurney M;

XX WPI; 2002-397167/43.

XX N-PSDB; ABL52487.

XX Human aspartyl protease 1 substrates useful in assays to detect aspartyl

XX protease activity, e.g. for the diagnosis of Alzheimer's disease.

CC The present invention describes a human aspartyl protease 1 (hu-Asp1)
CC substrate (I) which comprises a peptide of no more than 50 amino acids,
CC and which comprises the 8 amino acid sequence Gly-Leu-Ala-Leu-Ala-Leu-
CC Glu-Pro. Also described are: (1) a method (II) for assaying hu-Asp1
CC proteolytic activity, comprising: (a) contacting a hu-Asp1 protein with
CC (I) under acidic conditions; and (b) determining the level of hu-Asp1
CC proteolytic activity; (2) a purified polynucleotide (III) comprising a
CC nucleotide sequence that hybridizes under stringent conditions to the non
CC coding strand complementary to a defined 1804 nucleotide sequence (see
CC AB52456) where the nucleotide sequence encodes a transmembrane
CC proteolytic activity and lacks nucleotides encoding a transmembrane
CC domain; (3) a purified polynucleotide (III') comprising a sequence that
CC hybridizes under stringent conditions to (III) (the nucleotide sequence
CC encodes a polypeptide further lacking a pro-peptide domain corresponding
CC to amino acids 23-62 of hu-Asp1 (see AB878589)); (4) a vector (IV)
CC comprising (III) or (III') and (5) a host cell (V) transformed or
CC transfected with (III), (III') and/or (IV). The hu-Asp1 protease
CC substrate (I) may be used as an enzyme substrate in assays to detect
CC aspartyl protease activity, (II) and therefore diagnose diseases
CC associated with aberrant hu-Asp1 expression and activity such as
CC Alzheimer's disease. Hu-Asp1 has been localised to chromosome 21, while
CC hu-Asp2 has been localised to chromosome 11q23.3-24.1. The present
CC sequence represents human Asp-2(b) deltatm, which is given in an example
CC from the present invention
CC
XX
SQ Sequence 428 AA;

Query Match 100.0%; Score 16; DB 5; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TQHGIRLPRLSGGGA 16
|||
22 TQHGIRLPRLSGGGA 37

RESULT 9
ADJ94363
ID ADJ94363 standard; protein; 428 AA.

AC ADJ94363;

DT 03-JUN-2004 (first entry)

DE Human-pro-Asp-2(b) deltatm.

XX Human; enzyme; aspartyl protease; Asp-1; Asp-2(a); Asp-2(b);

XX beta secretase; amyloid protein precursor; APP; Alzheimer's disease;

XX nootropic; neuroprotective; amyloid beta; mutant; mutuin.

XX Homo sapiens.

XX Synthetic.

XX US6706485-B1.

XX 16-MAR-2004.

XX 12-APR-2000; 2000US-00548376.

XX 24-SEP-1998; 98US-0101594P.

XX 23-SEP-1999; 99US-00404133.

XX 23-SEP-1999; 99US-0155493P.

XX 13-OCT-1999; 99WO-US020881.

XX (PHAA) PHARMACIA & UPJOHN CO.

XX Gurney ME, Bienkowski MJ, Heinrikson RL, Parodi LA, Yan R;

XX WPI; 2004-236722/22.

XX N-PSDB; ADJ94362.

XX Identifying agents that modulate activity of Asp2 aspartyl protease

XX

PT useful for treating or preventing Alzheimer's disease involves comparing
PT APP processing activity of protease in presence and absence of test
PT agent.
XX
PS Example 10; SEQ ID NO 51; 109pp; English.
XX
CC The invention relates to identifying agents that modulate activity of
CC Asp2 (e.g. a beta-secretase, e.g. human Asp-2(b) appearing as ID 6,
CC encoded by ID 5) aspartyl protease, involves contacting Asp2 with amyloid
CC precursor protein (APP) in the presence and absence of a test agent,
CC where Asp2 is a recombinant polypeptide and processes APP into amyloid
CC beta, determining APP processing activity of Asp2 in presence and absence
CC of the test agent, and comparing the activities to identify agents that
CC modulate the activity of Asp2. Also disclosed are the cDNA and proteins
CC for human Asp-1 and Asp-2(a), mouse Asp-2(b), a vector comprising the
CC nucleic acid encoding Hu-Asp2 protease sequence, a host cell comprising
CC the vector and the method of producing Hu-Asp polypeptide, an isolated
CC antibody that specifically binds to Hu-Asp polypeptides, identifying a
CC cell that can be used to screen for inhibitors of beta secretase
CC activity, novel isoforms of amyloid protein precursor (APP), where the
CC last 2 carboxy terminus amino acids of that isoform are both lysine
CC residues (e.g. those designated APP695-KK or carrying the Swedish
CC mutation where KM at 595-596 is mutated to NL, designated e.g. APP695-SW
CC or APP695-SW-KK, or a V to F mutation at 642, e.g. APP695-VF, all useful
CC for assaying for beta secretase activity and screening for inhibitors of
CC beta-secretase) and polynucleotides that encode the APP proteins. The
CC method is useful for identifying agents that modulate the activity
CC (amyloid precursor protein processing activity) of Asp2 aspartyl
CC protease. Preferably, the method is useful for identifying agents that
CC inhibit Asp2 aspartyl protease activity. The inhibitors of amyloid
CC precursor protein processing, are useful for treating or preventing
CC Alzheimer's disease. The present sequence represents an aspartyl protease
CC mutant construct (e.g. lacking a transmembrane domain and/or including a
CC caspase cleavage site) used to investigate the cleavage activity of Asp2
CC proteins.
XX
SQ Sequence 428 AA;

Query Match 100.0%; Score 16; DB 8; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TQHGIRLPRLSGIGCA 16
|||
Db 22 TQHGIRLPRLSGIGCA 37

RESULT 10
AD050459
ID AD050459 standard; protein; 428 AA.
XX
AC AD050459;
XX
DT 29-JUL-2004 (first entry)
XX
DE Human Asp2(b) deltatm mutant protein.
XX
KM Aspartyl protease; Asp; beta secretase; amyloid precursor protein; APP;
KM Alzheimer's disease; gene therapy; human; mutant; mutein.
XX
OS Homo sapiens.
OS Synthetic.
OS
PN US6737510-B1.
PN
PD 18-MAY-2004.
PD
PF 12-APR-2000; 2000US-00548373.
PF
XX
XX 24-SEP-1998; 98US-0101594P.
PR 23-SEP-1999; 99US-00404133.
PR 23-SEP-1999; 99US-0155493P.
PR 23-SEP-1999; 99MO-US020881.

PR 13-OCT-1999; 99US-00416901.
XX
XX (PHAA) PHARMACIA & UPJOHN CO.
PA
XX
PI Gurney ME, Bienkowiaki MJ, Heinrikson RL, Parodi LA, Yan R;
XX
DR WPI; 2004-387112/36.
DR N-PSDB; AD050458.
XX
PT New Asp2 aspartyl protease protein comprising tripeptides DTG and DSG
PT involved in processing amyloid precursor protein into amyloid beta,
PT useful in preparing a composition for treating or preventing Alzheimer's
PT disease.
XX
PS Example 10; SEQ ID NO 51; 108pp; English.
XX
CC The invention relates to a method for identifying an agent that decreases
CC the protease activity of the aspartyl protease (Asp) polypeptide. It also
CC provides enzyme and enzymatic procedures for cleaving the beta secretase
CC cleavage site of the amyloid precursor protein (APP). The invention is
CC useful in preparing a composition for treating or preventing Alzheimer's
CC disease. It is also useful in gene therapy. The present sequence is human
CC Asp2(b) mutant protein. This sequence is used to illustrate the method of
CC the invention.
XX
SQ Sequence 428 AA;

Query Match 100.0%; Score 16; DB 8; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TQHGIRLPRLSGIGCA 16
|||
Db 22 TQHGIRLPRLSGIGCA 37

RESULT 11
ADR75372
ID ADR75372 standard; protein; 428 AA.
XX
AC ADR75372;
XX
DT 18-NOV-2004 (first entry)
XX
DE Human Asp2(b) deltatm mutant protein.
XX
KM Aspartyl protease; Asp; amyloid precursor protein; APP; amyloid beta;
KM chromosome identification; Alzheimer's disease; human; mutant.
XX
OS Homo sapiens.
OS Synthetic.
OS
PN US2004166507-A1.
PN
PD 26-AUG-2004.
PD
PF 29-AUG-2003; 2003US-00652045.
PF
XX
XX 24-SEP-1998; 98US-0101594P.
PR 23-SEP-1999; 99US-00404133.
PR 23-SEP-1999; 99US-0155493P.
PR 13-OCT-1999; 99US-00416901.
XX
PA (GURN/) GURNEY M E.
PA (BIEN/) BIENKOWAKI M J.
PA (HEIN/) HEINRIKSON R L.
PA (PARO/) PARODI L A.
PA (YANR/) YAN R.
XX
XX Gurney ME, Bienkowiaki MJ, Heinrikson RL, Parodi LA, Yan R;
PI WPI; 2004-624916/60.
XX N-PSDB; ADR75371.
DR

XX Novel purified/isolated polynucleotide encoding polypeptide having
PT aspartyl protease activity involved in processing amyloid precursor
PT protein into amyloid beta, useful in identifying agent decreasing
PT activity of aspartyl protease.

XX Example 10; SEQ ID NO 51; 107pp; English.

XX The invention relates to nucleic acid sequences encoding aspartyl
XX protease (Asp) polypeptides having aspartyl protease activity involved in
XX processing amyloid precursor protein (APP) into amyloid beta. The
XX invention also relates to a method for identifying an agent that
XX decreases the protease activity of the Asp. Asp DNA is useful in
XX chromosome identification as they can hybridise with a specific location
XX on a human chromosome and in identifying the relationship between genes
XX and diseases (particular gene responsible for causing diseases). It is
XX also useful for identifying candidates to modulate the progression of
XX Alzheimer's disease. Asp is useful in raising antibodies that are useful
XX in diagnostic assay for detecting Hu-Asp polypeptide expression. The
XX present sequence is the human Asp2(b)deltaTM mutant protein. This
XX sequence is used to illustrate the method of the invention.

XX Sequence 428 AA;

Query Match 100.0%; Score 16; DB 8; Length 428;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TQHGIRLPLRSLGGA 16
|||
DB 22 TQHGIRLPLRSLGGA 37

RESULT 12

ID ADC81561
ADG1561 standard; protein; 432 AA.

XX ADC81561;

XX 01-JAN-2004 (first entry)

XX Mature BACE p33K amino acid sequence SEQ ID NO:2.

XX human; BACE; modification; Pro33lys; pro-enzyme.

XX Synthetic.

XX Homo sapiens.

XX WO2003072733-A2.

XX 04-SEP-2003.

XX 21-FEB-2003; 2003WO-US005508.

XX 21-FEB-2002; 2002US-0358651P.

XX (PHAA) PHARMACIA & UPJOHN CO.

XX Chou K, Howe JM;

XX WPI: 2003-712719/67.

XX N-PSDB; ADC81562.

XX BACE polypeptides having Pro33lys modification, useful in determining
XX possible mutations, which will inhibit enzyme activity, and in
XX determining potential active site for target molecules.

XX Claim 10; SEQ ID NO 2; 38pp; English.

XX The present invention describes an isolated polypeptide (1) comprising or
XX consisting of a fully defined sequence of 432 amino acids (see ADC81561),
XX and comprising human BACE having the modification Pro33lys. Also
XX described: (1) a composition comprising an active human BACE enzyme

CC comprising the pro-enzyme sequence of BACE having the modification
CC Pro33lys; (2) an isolated polynucleotide comprising a sequence encoding
CC (1); (3) an isolated polynucleotide consisting or comprising of
CC nucleotides 70-1165 of a 1355-bp sequence (see ADC81562); (4) an
CC expression vector comprising the polynucleotide of (2), or a
CC polynucleotide sequence encoding a Pro33lys-BACE polypeptide, where the
CC expression vector can produce the Pro33lys-BACE polypeptide when present
CC in a compatible host cell, when cultured under conditions that allow
CC production; (5) a recombinant host cell comprising the expression vector;
CC and (6) producing a (active) Pro33lys-BACE polypeptide. The BACE
CC polypeptide having Pro33lys modification may be used in determining
CC possible mutations, which will inhibit enzyme activity, and in
CC determining potential active site for target molecules. The vector
CC comprising the BACE polynucleotide is useful for producing recombinant
CC BACE polypeptides having Pro33lys modification. The present sequence
CC represents the mature recombinant BACE p33K amino acid sequence used in
CC the exemplification of the present invention.

XX Sequence 432 AA;

Query Match 100.0%; Score 16; DB 7; Length 432;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TQHGIRLPLRSLGGA 16
|||
DB 1 TQHGIRLPLRSLGGA 16

RESULT 13

ID AAY88433
AAY88433 standard; protein; 433 AA.

XX AAY88433;

XX 12-SEP-2003 (revised)

XX 06-AUG-2003 (revised)

XX 03-AUG-2000 (first entry)

XX Human-pro-Asp-2(a)-deltaTM amino acid sequence.

XX Aspartyl protease; aspartase; amyloid precursor protein; APP; Asp 2;
XX Alzheimer's disease; beta secretase site; human-pro-Asp-2(a)-deltaTM.

XX Homo sapiens.

XX Enterobacteria phage T7.

XX WO200017369-A2.

XX 30-MAR-2000.

XX 23-SEP-1999; 99WO-US020881.

XX 24-SEP-1998; 98US-0101594P.

XX (PHAA) PHARMACIA & UPJOHN CO.

XX Gurney ME, Bienkowski MJ, Heinrikson RL, Parodi LA, Yan R;

XX WPI: 2000-303209/26.

XX N-PSDB; AAA15670.

XX New enzyme designated human aspartase useful in research into Alzheimer's
XX disease is capable of cleaving amyloid protein precursor at the beta
XX secretase site to produce amyloid beta peptide.

XX Example 9; Fig 8; 183pp; English.

XX This sequence represents a modified version of the human aspartase 2
XX (Asp2) amino acid sequence. The sequence is used in the bacterial
XX expression of human Asp2L. The invention relates to a protease (e.g.
XX Asp2) capable of cleaving the beta secretase site of amyloid precursor

CC protein (App). The protease contains a sequence encoding the amino acid
 CC sequence DTG and a sequence encoding DSG or DRG separated by 100-300
 CC amino acids. When mutated the App gene causes an autosomal dominant form
 CC of Alzheimer's disease. App localises to the cell surface membrane and
 CC have a single C-terminal transmembrane domain. Proteolytic processing of
 CC APP produces the amyloid beta protein, which is possibly very important
 CC in Alzheimer's disease. The invention includes a nucleotide sequence
 CC encoding the protease, a vector containing the nucleotide sequence, and a
 CC cell line comprising the vector. Methods for screening for inhibitors of
 CC beta secretase activity are also given in the invention. The human
 CC aspartase protein and nucleotide sequences and the methods for
 CC identifying inhibitors of the protease, are useful in the treatment of
 CC and research in to Alzheimer's disease. (Updated on 06-AUG-2003 to
 CC correct OS field.) (Updated on 12-SEP-2003 to standardise OS field)

CC Sequence 433 AA;

Query Match Best Local Similarity 100.0%; Score 16; DB 3; Length 433;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TOHGIRLPURSGIGA 16
 |||||
 Db 2 TOHGIRLPURSGIGA 17

RESULT 14

AAU07213
 XX AAU07213 standard; protein; 433 AA.

AC AAU07213;

DT 11-SEP-2003 (revised)

DT 24-OCT-2001 (first entry)

DE T7-human aspartyl protease 2a delcATM (low GC).

KW Human; aspartyl protease 1; Asp-1, neuroprotective;

KW aspartyl protease 2; Asp2; amyloid protein precursor; App;

KW beta-secretase; Alzheimer's disease.

OS Homo sapiens.

OS Enterobacteria phage T7.

XX MO200149097-A2.

XX 12-JUL-2001.

XX 09-MAY-2001; 2001WO-IB000797.

XX 09-MAY-2001; 2001WO-IB000797.

XX 09-MAY-2001; 2001WO-IB000797.

XX (BIEN/) BIENKOWSKI M J.

XX (GURN/) GURNEY M E.

XX (HEIN/) HEINRIKSON R L.

XX (PARO/) PARODI L A.

XX (YANR/) YAN R.

XX Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;

XX WPI; 2001-502548/55.

XX N-PSDB; AAS11713.

XX Novel purified polypeptide comprising fragment of mammalian aspartyl

XX protease 2, lacking Asp2 transmembrane domain and retaining beta

XX secretase activity of Asp2 useful for identifying inhibitors of Asp2

XX activity.

XX Example 9; Fig 8; 185pp; English.

XX The invention relates to a novel purified polypeptide comprising a

XX fragment of mammalian aspartyl protease 2 (Asp2) protein which lacks the

XX Asp2 transmembrane domain and the Asp2 protein, and where the polypeptide

CC and the fragment retain the beta-secretase activity of the mammalian Asp2
 CC protein. Also included is an isoform of amyloid protein precursor (APP)
 CC comprising the amino acid sequence of a App or its fragment containing an
 CC APP cleavage site recognizable by a mammalian beta-secretase, and further
 CC comprising two lysine residues at the carboxyl terminus of the amino acid
 CC sequence of the mammalian App or App fragment. The polypeptides are used
 CC for assaying for modulators of beta-secretase activity; identifying
 CC agents that inhibit the APP processing activity of human Asp2 aspartyl
 CC protease (Hu-Asp2); identifying agents that modulate the activity of Asp2
 CC ; and for reducing cellular production of amyloid beta (Abeta) from APP.
 CC Alzheimer's disease, for identifying modulators of amyloid-beta (Abeta)
 CC peptide production, and for use in designing therapeutics for the
 CC treatment or prevention of Alzheimer's disease. Probes and primers
 CC derived from Asp nucleic acid sequences are useful for detecting Hu-Asp
 CC nucleic acids in in vitro assays and in Northern and Southern blots. The
 CC present sequence represents the amino acid sequence of T7-human Asp-2a
 CC delta TM (low GC) construct which has a T7 tag, has the GC content of the
 CC 5' sequence reduced by site-directed mutagenesis, and lacks the
 CC transmembrane domain. This construct was used for bacterial expression
 CC and purification of human Asp2a. (Updated on 11-SEP-2003 to standardise
 CC OS field)

CC Sequence 433 AA;

Query Match Best Local Similarity 100.0%; Score 16; DB 4; Length 433;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TOHGIRLPURSGIGA 16
 |||||
 Db 2 TOHGIRLPURSGIGA 17

RESULT 15

AAE10640
 XX AAE10640 standard; protein; 433 AA.

AC AAE10640;

DT 10-DEC-2001 (first entry)

DE Human-pro-Asp 2(a) protein lacking TM domain.

KW Human; aspartyl protease 1; Asp1; amyloid precursor protein; App;

KW Alzheimer's disease; AD; dementia; neurofibrillary tangle; gliosis;

KW amyloid plaque; neuronal loss; proteolytic; neuroprotective;

KW Human-pro-Asp 2(a) protein.

OS Homo sapiens.

OS Synthetic.

XX GB2357767-A.

XX 04-JUL-2001.

XX 22-SEP-2000; 2000GB-00023315.

XX 23-SEP-1999; 99US-00404133.

XX 23-SEP-1999; 99US-0155493P.

XX 23-SEP-1999; 99WO-US0200881.

XX 13-OCT-1999; 99US-00416901.

XX 06-DEC-1999; 99US-0169232P.

XX (PHAA) PHARMACIA & UPJOHN CO.

XX Bienkowski MJ, Gurney M;

XX WPI; 2001-444206/48.

XX N-PSDB; AAD17876.

XX Polypeptide comprising fragments of human aspartyl protease with amyloid

XX precursor protein processing activity and alpha-secretase activity, for

PR identifying modulators useful in treating Alzheimer's disease.

XX
PS Example 9; Fig 8; 187pp; English.

XX The patent discloses human aspartyl protease 1 (hu-Asp1) or modified Asp1
CC proteins which lack transmembrane domain or amino terminal domain or
CC cytoplasmic domain and retains alpha-secretase activity and amyloid
CC protein precursor (APP) processing activity. The proteins of the
CC invention are useful for assaying hu-Asp1 alpha-secretase activity, which
CC in turn is useful for identifying modulators of hu-Asp1 alpha-secretase
CC activity, where modulators that increase hu-Asp1 alpha-secretase activity
CC are useful for treating Alzheimer's disease (AD) which causes progressive
CC dementia with consequent formation of amyloid plaques, neurofibrillary
CC tangles, gliosis and neuronal loss. Hu-Asp1 protease substrate is useful
CC for assaying hu-Asp1 proteolytic activity, by contacting hu-Asp1 protein
CC with the substrate under acidic conditions and determining the level of
CC hu-Asp1 proteolytic activity. The present sequence human-Pro-Asp 2(a)
CC protein lacking a transmembrane (TM) domain (low GC). This sequence is
CC generated from human Asp 2(a) protein by the deletion of its C-terminal
CC transmembrane domain and change of degenerate codon bases in 15 amino
CC acid positions from G/C to A/T to reduce the GC content

XX
SQ Sequence 433 AA;

Query Match 100.0%; Score 16; DB 4; Length 433;

Best Local Similarity 100.0%; Pred. No. 1,1e-07;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TOHGIRLPRLRSGLGGA 16
|||
Db 2 TOHGIRLPRLRSGLGGA 17

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Job time : 168 secs

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OM protein - protein search, using sw model

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(without alignments)
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	16	100.0	425	15	US-10-372-473-3
3	16	100.0	428	9	US-09-794-927-51
4	16	100.0	428	9	US-09-795-847-51
5	16	100.0	428	9	US-09-794-743-51
6	16	100.0	428	9	US-09-794-748-51
7	16	100.0	428	9	US-09-794-925-51
8	16	100.0	428	9	US-09-681-442-51
9	16	100.0	428	10	US-09-869-414-51
10	16	100.0	428	10	US-09-548-366-51
11	16	100.0	428	15	US-10-652-927-51

12	16	100.0	428	15	US-10-652-830-51	Sequence 51, Appl
13	16	100.0	428	16	US-10-652-045-51	Sequence 51, Appl
14	16	100.0	428	16	US-10-476-935-51	Sequence 51, Appl
15	16	100.0	428	17	US-10-477-076-51	Sequence 51, Appl
16	16	100.0	432	15	US-10-372-473-2	Sequence 2, Appl
17	16	100.0	433	9	US-09-794-927-26	Sequence 26, Appl
18	16	100.0	433	9	US-09-795-847-26	Sequence 26, Appl
19	16	100.0	433	9	US-09-794-743-26	Sequence 26, Appl
20	16	100.0	433	9	US-09-794-748-26	Sequence 26, Appl
21	16	100.0	433	9	US-09-794-925-26	Sequence 26, Appl
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23	16	100.0	433	10	US-09-869-414-26	Sequence 26, Appl
24	16	100.0	433	10	US-09-548-366-26	Sequence 26, Appl
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28	16	100.0	433	16	US-10-476-935-26	Sequence 26, Appl
29	16	100.0	433	17	US-10-940-867-26	Sequence 26, Appl
30	16	100.0	433	17	US-10-726-967A-78	Sequence 78, Appl
31	16	100.0	433	17	US-10-726-967A-81	Sequence 81, Appl
32	16	100.0	433	17	US-10-726-967A-84	Sequence 84, Appl
33	16	100.0	433	17	US-10-477-076-26	Sequence 26, Appl
34	16	100.0	434	9	US-09-794-927-53	Sequence 53, Appl
35	16	100.0	434	9	US-09-795-847-53	Sequence 53, Appl
36	16	100.0	434	9	US-09-794-743-53	Sequence 53, Appl
37	16	100.0	434	9	US-09-794-748-53	Sequence 53, Appl
38	16	100.0	434	9	US-09-794-925-53	Sequence 53, Appl
39	16	100.0	434	9	US-09-681-442-53	Sequence 53, Appl
40	16	100.0	434	10	US-09-869-414-53	Sequence 53, Appl
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43	16	100.0	434	15	US-10-652-830-53	Sequence 53, Appl
44	16	100.0	434	16	US-10-652-045-53	Sequence 53, Appl
45	16	100.0	434	16	US-10-476-935-53	Sequence 53, Appl

ALIGNMENTS

RESULT 1
US-10-726-967A-3
; Sequence 3, Application US/10726967A
; Publication No. US20050074456A1
; GENERAL INFORMATION:
; APPLICANT: Ballinger, Marcus
; TITLE OF INVENTION: Construct for Homogenously Processed Preparations of Beta Site
; FILE REFERENCE: 2004345-0021
; CURRENT APPLICATION NUMBER: US/10/726,967A
; CURRENT FILING DATE: 2003-12-02
; NUMBER OF SEQ ID NOS: 110
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 3
; LENGTH: 16
; TYPE: PRT
; ORGANISM: Artificial
; FEATURE:
; OTHER INFORMATION: Residues 22-37 of human BACE1 preprosequence
US-10-726-967A-3

Query Match 100.0%; Score 16; DB 17; Length 16;
Best Local Similarity 100.0%; Pred. No. 1.7e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 1 TQHGRLPLRSGIGGA 16
RESULT 2
US-10-372-473-3
; Sequence 3, Application US/10372473
; Publication No. US20040005691A1

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; GENERAL INFORMATION:
; APPLICANT: Chou, Kuo-Chen
; TITLE OF INVENTION: Modified BACE
; FILE REFERENCE: MBAB 01-1766-A
; CURRENT APPLICATION NUMBER: US/10/372,473
; NUMBER OF SEQ ID NOS: 24
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO: 3
; LENGTH: 425
; TYPE: PRT
; ORGANISM: Homo sapiens
; NAME/KEY: MISC FEATURE
; OTHER INFORMATION: Human beta-secretase zymogen.
US-10-372-473-3

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Query Match          100.0%; Score 16; DB 15; Length 425;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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        1 TOHGIRLPLRSGIGA 16

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RESULT 3
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; Sequence 51, Application US/09794927
; Patent No. US20010016324A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; FILE REFERENCE: 28341/6280FG
; CURRENT APPLICATION NUMBER: US/09/794,927
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-794-927-51

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Query Match          100.0%; Score 16; DB 9; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY      1 TOHGIRLPLRSGIGA 16
        |||||
        22 TOHGIRLPLRSGIGA 37

```

```

RESULT 4
US-09-795-847-51
; Sequence 51, Application US/09795847
; Patent No. US20010018208A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; FILE REFERENCE: 28341/6280DE
; CURRENT APPLICATION NUMBER: US/09/795,847
; PRIOR FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-795-847-51

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```

Query Match          100.0%; Score 16; DB 9; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY      1 TOHGIRLPLRSGIGA 16
        |||||
        22 TOHGIRLPLRSGIGA 37

```

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RESULT 5
US-09-794-743-51
; Sequence 51, Application US/09794743
; Patent No. US20010021391A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; FILE REFERENCE: 28341/6280BC
; CURRENT APPLICATION NUMBER: US/09/794,743
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73

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; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-794-743-51

Query Match          100.0%; Score 16; DB 9; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGGGA 16
        |||||||
        22 TOHGIRLPLRSGGGA 37

DB

RESULT 6
US-09-794-748-51
; Sequence 51, Application US/09794748
; Patent No. US20020037315A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280JL
; CURRENT APPLICATION NUMBER: US/09/794,748
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-794-748-51

Query Match          100.0%; Score 16; DB 9; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGGGA 16
        |||||||
        22 TOHGIRLPLRSGGGA 37

DB

RESULT 7
US-09-794-925-51
; Sequence 51, Application US/09794925
; Patent No. US20020064819A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280HI
; CURRENT APPLICATION NUMBER: US/09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-681-442-51

Query Match          100.0%; Score 16; DB 9; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 TOHGIRLPLRSGGGA 16
        |||||||
        22 TOHGIRLPLRSGGGA 37

DB

RESULT 8
US-09-681-442-51
; Sequence 51, Application US/09681442
; Patent No. US20020081634A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280FG
; CURRENT APPLICATION NUMBER: US/09/681,442
; PRIOR FILING DATE: 2001-04-05
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-681-442-51
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Query Match 100.0%; Score 16; DB 9; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 9

US-09-869-414-51
; Sequence 51, Application US/09869414
; Publication No. US20030077226A1
; GENERAL INFORMATION:
; APPLICANT: Beinikowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M
; CURRENT APPLICATION NUMBER: US/09/869,414
; PRIOR FILING DATE: 2001-06-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-869-414-51

Query Match 100.0%; Score 16; DB 10; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 10

US-09-548-366-51
; Sequence 51, Application US/09548366
; Publication No. US20030104365A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Beinikowski, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES THEREFOR
; FILE REFERENCE: 28341/6280A
; CURRENT APPLICATION NUMBER: US/09/548,366
; PRIOR FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24

; NUMBER OF SEQ ID NOS: 65
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
; OTHER INFORMATION: delta TM
US-09-548-366-51

Query Match 100.0%; Score 16; DB 10; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 11

US-10-652-927-51
; Sequence 51, Application US/10652927
; Publication No. US20040043408A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280N3
; CURRENT APPLICATION NUMBER: US/10/652,927
; PRIOR FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: 09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 51
; LENGTH: 428
; TYPE: PRT
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-10-652-927-51

Query Match 100.0%; Score 16; DB 15; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 12

US-10-652-830-51
; Sequence 51, Application US/10652830
; Publication No. US20040048303A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280N1
; CURRENT APPLICATION NUMBER: US/10/652,830
; PRIOR FILING DATE: 2003-08-29

PRIOR APPLICATION NUMBER: 09/794,925
PRIOR FILING DATE: 2001-02-27
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 74
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-10-652-830-51

Query Match 100.0%; Score 16; DB 15; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 13
US-10-652-045-51
Sequence 51, Application US/10652045
Publication No. US2004016507A1
GENERAL INFORMATION:
APPLICANT: Gurney et al.
TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
FILE REFERENCE: 29915/6280N2
CURRENT APPLICATION NUMBER: US/10/652,045
CURRENT FILING DATE: 2003-08-29
PRIOR APPLICATION NUMBER: 09/794,925
PRIOR FILING DATE: 2001-02-27
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 74
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-10-652-045-51

Query Match 100.0%; Score 16; DB 16; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 14
US-10-476-935-51
Sequence 51, Application US/10476935
Publication No. US20040234976A1
GENERAL INFORMATION:
APPLICANT: Beinkowski et al.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
FILE REFERENCE: 28341/6280M1
CURRENT APPLICATION NUMBER: US/10/476,935
CURRENT FILING DATE: 2003-11-06
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
US-10-476-935-51

Query Match 100.0%; Score 16; DB 16; Length 428;
Best Local Similarity 100.0%; Pred. No. 2.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 TOHGIRLPLRSGGGA 16
|||
Db 22 TOHGIRLPLRSGGGA 37

RESULT 15
US-10-477-076-51
Sequence 51, Application US/10477076
Publication No. US2005008022A1
GENERAL INFORMATION:
APPLICANT: Beinkowski et al.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
FILE REFERENCE: 28341/6280M2
CURRENT APPLICATION NUMBER: US/10/477,076
CURRENT FILING DATE: 2003-11-06
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 51
LENGTH: 428
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
US-10-477-076-51

Query Match 100.0%; Score 16; DB 17; Length 428;
 Best Local Similarity 100.0%; Pred. No. 2.6e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TQHGIRLPLRSGLGGA 16
 |||||
 Db 22 TQHGIRLPLRSGLGGA 37

Search completed: July 26, 2005, 16:41:29
 Job time : 154 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: July 26, 2005, 16:28:04 ; Search time 39 Seconds
(without alignments)
39.474 Million cell updates/sec

Title: US-10-726-967A-3

Perfect score: 16

Sequence: 1 TQHGIRLPLRSGLGGA 16

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 283416 segs, 96216763 residues

Word size : 0

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 45 summaries

Database : PIR 79:*

1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	16	100.0	501	2	A59090	aspartic proteinase
2	7	43.8	125	2	E81814	hypothetical prote
3	7	43.8	239	2	AG0420	phosphonates trans
4	7	43.8	259	2	A75547	hypothetical prote
5	7	43.8	352	2	D64966	membrane protein Y
6	6	37.5	157	2	AG2675	hypothetical prote
7	6	37.5	169	2	E90983	probable GDP-L-fuc
8	6	37.5	169	2	H85828	GDP-mannose mannos
9	6	37.5	225	2	D81813	hypothetical prote
10	6	37.5	237	2	T35108	hypothetical prote
11	6	37.5	241	2	T26674	hypothetical prote
12	6	37.5	246	2	AG3644	flagellar biosynth
13	6	37.5	252	2	AE3631	nitrous-oxide redu
14	6	37.5	274	2	D97653	hypothetical prote
15	6	37.5	274	2	AB2877	conserved hypothet
16	6	37.5	284	2	B41224	homeotic protein p
17	6	37.5	292	2	C83520	dihydrodipicolinat
18	6	37.5	299	2	B95149	heat shock protein
19	6	37.5	302	2	A99017	heat shock protein
20	6	37.5	325	2	A72724	hypothetical prote
21	6	37.5	338	2	AE3334	metal chelate tran
22	6	37.5	352	2	T02875	ribosomal protein
23	6	37.5	352	2	D85826	probable transpor
24	6	37.5	352	2	G90980	probable transpor
25	6	37.5	379	2	A57477	potassium channel
26	6	37.5	390	2	AF3425	oxidoreductase (EC
27	6	37.5	399	2	AC2785	MFS permease (drug
28	6	37.5	399	2	D97564	hypothetical prote
29	6	37.5	411	2	T34585	probable secreted

30	6	37.5	416	2	T32458	hypothetical prote
31	6	37.5	423	2	A45363	somatoliberein rece
32	6	37.5	430	2	A84165	UDP-glucose dehydr
33	6	37.5	430	2	D96536	hypothetical prote
34	6	37.5	437	2	D80151	conserved hypothet
35	6	37.5	439	2	D70954	hypothetical glyci
36	6	37.5	444	2	B65045	hypothetical prote
37	6	37.5	444	2	C85913	hypothetical prote
38	6	37.5	444	2	A91069	hypothetical prote
39	6	37.5	451	2	I46586	growth hormone-rel
40	6	37.5	454	2	H83377	probable transpor
41	6	37.5	463	2	G83175	probable metallo-o
42	6	37.5	464	2	S29754	growth hormone-rel
43	6	37.5	504	2	AD3629	vdcc protein (limp
44	6	37.5	518	2	S75811	gamma-glutamyltran
45	6	37.5	525	2	AF2950	GDH family prote

ALIGNMENTS

RESULT 1
A59090
aspartic proteinase (EC 3.4.23.-) BACE precursor - human
N:Alternate names: beta-secretase; beta-site APP cleaving enzyme
C:Species: Homo sapiens (man)
C>Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 09-Jul-2004
C:Accession: A59090
R:Vassar, R.; Bennett, B.D.; Babu-Khan, S.; Kahn, S.; Mendiaz, E.A.; Denis, P.; Teplow, M.A.; Biere, A.L.; Curran, E.; Burgess, T.; Louis, J.C.; Collins, F.; Treanor, J.; Rogers, Science 286, 735-741, 1999
A>Title: beta-Secretase cleavage of Alzheimer's amyloid precursor protein by the transmembrane
A:Reference number: A59090 ; MUID:20002972 ; PMID:10531052
A>Note: submitted to Genbank, September 1999
A:Accession: A59090
A>Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-501 <VAS>
A:Cross-references: UNIPROT:P56817; GB:AF190725; NID:g6118538; PIDN:AAF04142.1; PID:g611
C:Genetics:
A:Gene: BACE
C:Superfamily: beta-secretase
C:Keywords: Alzheimer's disease; aspartic proteinase; brain; glycoprotein; hydrolase; pr
F:1-21/Domain: signal sequence #status predicted <SIG>
F:22-45/Domain: propeptide #status predicted <PRO>
F:46-501/Product: acid proteinase BACE #status predicted <MAT>
F:461-477/Domain: transmembrane #status predicted <TRN>
F:33,289/Active site: Asp #status predicted
F:153,172,223,354/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:330-380/Disulfide bonds: #status predicted

Query Match 100.0%; Score 16; DB 2; Length 501;
Best Local Similarity 100.0%; Pred. No. 1.8e-09;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TQHGIRLPLRSGLGGA 16
DB 22 TQHGIRLPLRSGLGGA 37

RESULT 2
E81814
hypothetical protein NMA1874 [imported] - Neisseria meningitidis (strain Z2491 serogrou
C:Species: Neisseria meningitidis
C>Date: 05-May-2000 #sequence_revision 05-May-2000 #text_change 09-Jul-2004
C:Accession: E81814
R:Parkhill, J.; Achtman, M.; James, K.D.; Bentley, S.D.; Churruarin, C.; Klee, S.R.; Morel
; Holroyd, S.; Jagels, K.; Leather, S.; Moule, S.; Mungall, K.; Quail, M.A.; Rajadream,
Nature 404, 502-506, 2000
A>Title: Complete DNA sequence of a serogroup A strain of Neisseria meningitidis Z2491.
A:Reference number: AB1775; MUID:20222556; PMID:10761919
A:Accession: E81814
A>Status: preliminary

A:Molecule type: DNA
A:Residues: 1-125 <PAR>
A:Cross-references: UNIPROT:Q9JTC5; GB:AL162757; GB:AL157959; NID:97380371; PIDN:CAB8509
A:Experimental source: serogroup A, strain 22491
C:Genetics:
A:Gene: NMA1874
C:Superfamily: Neisseria meningitidis hypothetical protein NMA1874

Query Match 43.8%; Score 7; DB 2; Length 125;
Best Local Similarity 100.0%; Pred. No. 1.8;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 9 LRSGLG 15
DB 109 LRSGLG 115

RESULT 3
AG0420 phosphonates transport ATP-binding protein phnL [imported] - Yersinia pestis (strain CO3
C:Species: Yersinia pestis
C>Date: 02-Nov-2001 #sequence_revision 02-Nov-2001 #text_change 09-Jul-2004
C:Accession: AG0420
R:Barhill, J.; Wren, B.W.; Thomson, N.R.; Tibball, R.W.; Holden, M.T.G.; Prentice, M.B.
deno-Farraga, A.M.; Chillingworth, T.; Cronin, A.; Davies, R.M.; Davis, P.; Dougan, G.;
11, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; Whitehead, S.; Barrett,
Nature 413, 523-527, 2001
A:Title: Genome sequence of Yersinia pestis, the causative agent of plague.
A:Reference number: AB0001; MUID:21470413; PMID:11586360
A:Accession: AG0420
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-239 <KUR>
A:Cross-references: UNIPROT:Q8ZBF4; GB:AL590842; PIDN:CMC92691.1; PID:G15981386; GSPDB:C
A:Genetics:
C:Superfamily: short-chain ATP-binding cassette proteins; ATP-binding cassette homology

Query Match 43.8%; Score 7; DB 2; Length 239;
Best Local Similarity 100.0%; Pred. No. 3.2;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 OHGRLP 8
DB 20 OHGRLP 26

RESULT 4
A75547 hypothetical protein - Deinococcus radiodurans (strain R1)
C:Species: Deinococcus radiodurans
C>Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 09-Jul-2004
C:Accession: A75547
R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.;
S.; Smith, H.O.; Venter, J.C.; Lam, P.; McDonald, L.; Uterback, T.; Zalewski, C.; Ma
Science 286, 1571-1577, 1999
A:Title: Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1.
A:Reference number: A75540; MUID:20036896; PMID:10567266
A:Accession: A75547
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-259 <WHI>
A:Cross-references: UNIPROT:Q9RXU2; GB:AE001883; GB:AE000513; NID:96457878; PIDN:AAF0980
A:Experimental source: strain R1
C:Genetics:
A:Gene: DR0214
A:Map position: 1
C:Superfamily: Deinococcus radiodurans hypothetical protein DR0214

Query Match 43.8%; Score 7; DB 2; Length 259;
Best Local Similarity 100.0%; Pred. No. 3.5;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 HGIRLPL 9
DB 190 HGIRLPL 196

RESULT 5
D64966 membrane protein yeeB - Escherichia coli (strain K-12)
C:Species: Escherichia coli
C>Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 09-Jul-2004
C:Accession: D64966
R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; Co.
A.; Rose, D.U.; Mau, B.; Shao, Y.
Science 277, 1453-1462, 1997
A:Title: The complete genome sequence of Escherichia coli K-12.
A:Reference number: A64720; MUID:97426617; PMID:9278503
A:Accession: D64966
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-352 <BLAT>
A:Cross-references: UNIPROT:P33015; GB:AE000292; GB:U00096; NID:91788310; PIDN:AACT5074.1
A:Experimental source: strain K-12, substrain MG1655
C:Genetics:
A:Gene: yeeB
C:Keywords: transmembrane protein
F:2-18/Domain: transmembrane #status predicted <TM1>
F:44-60/Domain: transmembrane #status predicted <TM2>
F:75-91/Domain: transmembrane #status predicted <TM3>
F:106-122/Domain: transmembrane #status predicted <TM4>
F:153-169/Domain: transmembrane #status predicted <TM5>
F:201-217/Domain: transmembrane #status predicted <TM6>
F:250-266/Domain: transmembrane #status predicted <TM7>
F:320-336/Domain: transmembrane #status predicted <TM8>

Query Match 43.8%; Score 7; DB 2; Length 352;
Best Local Similarity 100.0%; Pred. No. 4.6;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 LRSGLG 15
DB 283 LRSGLG 289

RESULT 6
AG2675 hypothetical protein Atu0805 [imported] - Agrobacterium tumefaciens (strain C58, Dupont)
C:Species: Agrobacterium tumefaciens
C>Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 09-Jul-2004
C:Accession: AG2675
R:Wood, D.W.; Setubal, J.C.; Kaul, R.; Monke, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo, L.
erge, G.; Gilliet, W.; Grant, C.; Guenicher, D.; Kutayavin, T.; Levy, R.; Li, M.; McCelli,
Science 294, 2317-2323, 2001
A:Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm,
ster, E.W.
A:Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
A:Reference number: AB2577; MUID:21608550; PMID:11743193
A:Accession: AG2675
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-157 <KUR>
A:Cross-references: UNIPROT:Q8UH78; GB:AE006688; PIDN:AL41821.1; PID:917739178; GSPDB:G
A:Experimental source: strain C58 (Dupont)
C:Genetics:
A:Gene: Atu0805
A:Map position: circular chromosome

Query Match 37.5%; Score 6; DB 2; Length 157;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 8 LRSGL 13

Db 81 PLRSL 86

RESULT 7

E90983
probable GDP-L-fucose pathway enzyme [imported] - Escherichia coli (strain O157:H7, sub
C:Species: Escherichia coli
C:Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 16-Aug-2004
C:Accession: E90983
R:Hayashi, T.; Makino, K.; Onishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.
gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
DNA Res. 8, 11-22, 2001
A:Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and genc
A:Reference number: A99629; MUID:21156231; PMID:11258796
A:Accession: E90983
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-169 <HAY>
A:Cross-references: UNIPROT:O85341; GB:BA000007; PIDN:BA36260.1; PID:g13362305; GSPDB:C
A:Experimental source: strain O157:H7, substrain RMD 0509952
C:Genetics:
A:Gene: ECe2837
C:Superfamily: mut domain homology

Query Match 37.5%; Score 6; DB 2; Length 169;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GIRLPL 9
Db 71 GIRLPL 76

RESULT 8

H85828
GDP-mannose mannosylhydrolase [imported] - Escherichia coli (strain O157:H7, substrain H
C:Species: Escherichia coli
C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 16-Aug-2004
C:Accession: H85828
R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew
Miller, L.; Grobeck, E.J.; Davis, N.W.; Llin, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
Nature 409, 529-533, 2001
A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A:Reference number: AB5480; MUID:21074935; PMID:11206551
A:Accession: H85828
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-169 <STO>
A:Cross-references: UNIPROT:O85341; GB:AE005174; NID:g12516220; PIDN:AA57092.1; GSPDB:C
A:Experimental source: strain O157:H7, substrain EDL933
C:Genetics:
A:Gene: wbdQ
C:Superfamily: mut domain homology

Query Match 37.5%; Score 6; DB 2; Length 169;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GIRLPL 9
Db 71 GIRLPL 76

RESULT 9

D81813
hypothetical protein NMA1865 [imported] - Neisseria meningitidis (strain Z2491 serogroup
C:Species: Neisseria meningitidis
C:Date: 05-May-2000 #sequence_revision 05-May-2000 #text_change 09-Jul-2004
C:Accession: D81813
R:Parkhill, J.; Achtman, M.; James, K.D.; Bentley, S.D.; Churcher, C.; Klee, S.R.; Morel
; Holroyd, S.; Jagers, K.; Leather, S.; Moule, S.; Mungall, K.; Quail, M.A.; Rajandream,
Nature 404, 502-506, 2000

A:Title: Complete DNA sequence of a serogroup A strain of Neisseria meningitidis Z2491.
A:Reference number: A81775; MUID:20222556; PMID:10761919
A:Accession: D81813
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-225 <PAR>
A:Cross-references: UNIPROT:Q9J968; GB:AL162757; GB:AL157959; NID:g7380371; PIDN:CAB8508
A:Experimental source: serogroup A, strain Z2491
C:Genetics:
A:Gene: NMA1865

Query Match 37.5%; Score 6; DB 2; Length 225;
Best Local Similarity 100.0%; Pred. No. 35;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GIRLPL 9
Db 197 GIRLPL 202

RESULT 10

T35108
hypothetical protein SC4H2.09 SC4H2.09 - Streptomyces coelicolor
C:Species: Streptomyces coelicolor
C:Date: 05-Nov-1999 #sequence_revision 05-Nov-1999 #text_change 09-Jul-2004
C:Accession: T35108
R:Seeger, K.J.; Harris, D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
submitted to the EMBL Data Library, March 1998
A:Reference number: Z21568
A:Accession: T35108
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-237 <SEE>
A:Cross-references: UNIPROT:O69964; EMBL:AL022268; PIDN:CAA18325.1; GSPDB:GN00070; SCOE
A:Experimental source: strain A3(2)
C:Genetics:
A:Gene: SCOE:SC4H2.09
C:Superfamily: Streptomyces coelicolor hypothetical protein SC4H2.09

Query Match 37.5%; Score 6; DB 2; Length 237;
Best Local Similarity 100.0%; Pred. No. 37;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 SGLGGA 16
Db 214 SGLGGA 219

RESULT 11

T26676
hypothetical protein Y38F1A.1 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
C:Accession: T26676
R:Wallis, J.
submitted to the EMBL Data Library, October 1998
A:Reference number: Z20253
A:Accession: T26676
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-241 <WIL>
A:Cross-references: UNIPROT:Q9XWMS; EMBL:AL032639; PIDN:CAA21628.1; GSPDB:GN00020; CESP:
A:Experimental source: clone Y38F1A
C:Genetics:
A:Gene: CESP:Y38F1A.1
A:Map position: 2
A:Introns: 37/3; 76/2; 130/3; 202/3

Query Match 37.5%; Score 6; DB 2; Length 241;
Best Local Similarity 100.0%; Pred. No. 37;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 SGLGGA 16

Db 101 SGLGGA 106

RESULT 12

AG3644

A:Title: flagellar biosynthetic protein flpP [imported] - Brucella melitensis (strain 16M)

C:Species: Brucella melitensis

C:Date: 01-Feb-2002 #sequence_revision 01-Feb-2002 #text_change 09-Jul-2004

C:Accession: AG3644

R:DelVecchio, V.G.; Kapral, V.; Redkar, R.U.; Patra, G.; Mujer, C.; Los, T.; Ivanova,

Proc. Natl. Acad. Sci. U.S.A. 99, 443-448, 2002

A:Title: The genome sequence of the facultative intracellular pathogen Brucella melitensis

A:Reference number: AD3252; PMID:11756688

A:Accession: AG3644

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-246 <KUR>

A:Cross-references: UNIPROT:Q8VB21; GB:AE008918; PIDN:AAL54322.1; PID:G17985302; GSPDB:C

A:Experimental source: strain 16M

C:Genetics:

A:Gene: BME11080

A:Map position: 11

C:Superfamily: flagellar biosynthetic protein flpP

Query Match 37.5%; Score 6; DB 2; Length 246;

Best Local Similarity 100.0%; Pred. No. 38;

Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 RSGGGA 14

Db 75 RSGGGA 80

RESULT 13

AE3631

A:Title: nitrous-oxide reductase (EC 1.7.99.6) [imported] - Brucella melitensis (strain 16M)

C:Species: Brucella melitensis

C:Date: 01-Feb-2002 #sequence_revision 01-Feb-2002 #text_change 09-Jul-2004

C:Accession: AE3631

R:DelVecchio, V.G.; Kapral, V.; Redkar, R.U.; Patra, G.; Mujer, C.; Los, T.; Ivanova,

Proc. Natl. Acad. Sci. U.S.A. 99, 443-448, 2002

A:Title: The genome sequence of the facultative intracellular pathogen Brucella melitensis

A:Reference number: AD3252; PMID:11756688

A:Accession: AE3631

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-252 <KUR>

A:Cross-references: UNIPROT:Q8VB21; GB:AE008918; PIDN:AAL54216.1; PID:G17985186; GSPDB:C

A:Experimental source: strain 16M

C:Genetics:

A:Gene: BME10974

A:Map position: 11

C:Keywords: oxidoreductase

Query Match 37.5%; Score 6; DB 2; Length 252;

Best Local Similarity 100.0%; Pred. No. 39;

Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 SGLGGA 16

Db 28 SGLGGA 33

RESULT 14

D97653

A:Title: hypothetical protein AGR_C_4436 [imported] - Agrobacterium tumefaciens (strain C58, Cere

C:Species: Agrobacterium tumefaciens

C:Date: 30-Sep-2001 #sequence_revision 30-Sep-2001 #text_change 09-Jul-2004

C:Accession: D97653

R:Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Qurollo, B.; Goldman,

A.; Liu, F.; Mollam, C.; Allinger, M.; Dougherty, D.; Scott, C.; Lappas, C.; Markelz, B.; Science 294, 2323-2328, 2001

A:Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tumefaciens

A:Reference number: A97359; MUID:21608551; PMID:11743194

A:Accession: D97653

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-274 <KUR>

A:Cross-references: UNIPROT:Q8UCN9; GB:AE007869; PIDN:AAK8181.1; PID:G15157627; GSPDB:G

C:Genetics:

A:Gene: AGR_C_4436

A:Map position: circular chromosome

Query Match 37.5%; Score 6; DB 2; Length 274;

Best Local Similarity 100.0%; Pred. No. 42;

Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 10 RSGGGA 15

Db 249 RSGGGA 254

RESULT 15

AB2877

A:Title: conserved hypothetical protein Atu2444 [imported] - Agrobacterium tumefaciens (strain C58)

C:Species: Agrobacterium tumefaciens

C:Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 09-Jul-2004

C:Accession: AB2877

R:Wood, D.W.; Setubal, J.C.; Kaul, R.; Monke, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo, L.

erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutayvin, T.; Levy, R.; Li, M.; McClellan,

Science 294, 2317-2323, 2001

A:Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm,

ster, E.W.

A:Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.

A:Reference number: AB2577; MUID:21608550; PMID:11743193

A:Accession: AB2877

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-274 <KUR>

A:Cross-references: UNIPROT:Q8UCN9; GB:AE008688; PIDN:AAL43432.1; PID:G17740934; GSPDB:G

A:Experimental source: strain C58 (Dupont)

C:Genetics:

A:Gene: Atu2444

A:Map position: circular chromosome

Query Match 37.5%; Score 6; DB 2; Length 274;

Best Local Similarity 100.0%; Pred. No. 42;

Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 10 RSGGGA 15

Db 249 RSGGGA 254

Search completed: July 26, 2005, 16:38:03
Job time : 41 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: July 26, 2005, 16:27:19 ; Search time 173 Seconds
(without alignments)
47.360 Million cell updates/sec

Title: US-10-726-967A-3
Perfect score: 16
Sequence: 1 TQHGIRLPURSGLGGA 16

Scoring table: OLIGO
Gapop 60.0 , Gapept 60.0

Searched: 1612378 seqs, 512079187 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database : Uniprot 03: *
1: uniprot_sprot: *
2: uniprot_trembl: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	127	2	Q76KP0
2	16	100.0	501	1	BAEI_HUMAN
3	16	100.0	501	2	Q81YC8
4	10	62.5	467	2	Q8C4F4
5	10	62.5	501	1	BAEI_MOUSE
6	10	62.5	501	1	BAEI_RAT
7	10	62.5	501	2	Q8B0V4
8	10	62.5	501	2	Q8C7R1
9	8	50.0	814	2	Q7S2I8
10	7	43.8	97	2	Q9JPF6
11	7	43.8	125	2	Q9UTC5
12	7	43.8	181	2	Q7NTR4
13	7	43.8	239	2	Q66F29
14	7	43.8	239	2	Q8ZBF4
15	7	43.8	259	2	Q9RXU2
16	7	43.8	272	2	Q67QZ3
17	7	43.8	293	2	Q74X41
18	7	43.8	299	2	Q9JHS7
19	7	43.8	304	2	Q7SFA9
20	7	43.8	329	2	Q7NTO7
21	7	43.8	350	2	Q93F54
22	7	43.8	352	1	YEEB_ECOLI
23	7	43.8	352	2	Q93F48
24	7	43.8	352	2	Q93F55
25	7	43.8	352	2	Q93F57
26	7	43.8	352	2	Q93F60
27	7	43.8	352	2	Q93R10
28	7	43.8	353	2	Q93F51
29	7	43.8	354	2	Q93F52
30	7	43.8	354	2	Q93F61
31	7	43.8	457	2	Q886W9

32	7	43.8	460	2	Q9X3V2	Q9X3V2 pseudomonas
33	7	43.8	468	2	Q88P19	Q88P19 pseudomonas
34	7	43.8	629	2	Q67Q15	Q67Q15 symbiodace
35	7	43.8	889	2	Q42723	Q42723 emericella
36	7	43.8	911	1	CAFA_MOUSE	Q9QWFO mus musculus
37	7	43.8	1104	2	Q9F060	Q9F060 oryza sativ
38	6	37.5	55	2	Q8BK1	Q8BK1 xanthomonas
39	6	37.5	55	2	Q7UVPI	Q7UVPI rhodospirillum rubrum
40	6	37.5	61	2	Q7J3N8	Q7J3N8 neurospora
41	6	37.5	68	2	Q6J3X9	Q6J3X9 burkholderia
42	6	37.5	77	2	Q7AET6	Q7AET6 geobacter
43	6	37.5	99	2	Q949C5	Q949C5 oryza sativ
44	6	37.5	99	2	Q6H7G5	Q6H7G5 oryza sativ
45	6	37.5	100	2	Q6ZB08	Q6ZB08 oryza sativ

ALIGNMENTS

RESULT 1
ID Q76KP0 PRELIMINARY; PRT; 127 AA.
AC Q76KP0;
DT 05-JUL-2004 (TREMBLrel. 27, Created)
DT 05-JUL-2004 (TREMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TREMBLrel. 27, Last annotation update)
DE Beta-site APP cleaving enzyme isoform I-127.
GN Name=BACE;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_Taxid=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Tanahashi H.;
RL Submitted (Aug-2002) to the EMBL/GenBank/DBJ databases.
CC -1 SIMILARITY: Belongs to the EMBL/GenBank/DBJ databases.
DR EMBL; AB089958; BAC81826.1; -
DR HSSP; P00797; 1BBS.
DR GO; GO:0003049; F.aspartic-type signal peptidase activity; IEA.
DR GO; GO:0005508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR001461; Peptidase_A1.
DR InterPro; IPR009119; Pept_A1_BACE.
DR InterPro; IPR009120; Pept_A1_BACE.
DR InterPro; IPR009007; Pept_Aspartic.
DR InterPro; IPR001969; Pept_Asp_AS.
DR Pfam; PF00026; Asp; 1.
DR PRINTS; PRO1815; BACEFAMILY.
DR PROSITE; PS00141; ASP_PROTEASE; 1.
KW Aspartyl protease; Hydrolyase; Protease.
SQ SEQUENCE 127 AA; 1393 MW; C657354CB872DC4 CRC64;

Query Match 100.0%; Score 16; DB 2; Length 127;
Best Local Similarity 100.0%; Pred. No. 1.3e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TQHGIRLPURSGLGGA 16
Db 22 TQHGIRLPURSGLGGA 37

RESULT 2
ID BAEI_HUMAN STANDARD; PRT; 501 AA.
AC P56817; Q9BYB9; Q9BYC0; Q9BYC1; Q9JUT5;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Beta-secretase 1 precursor (EC 3.4.23.46) (Beta-site APP cleaving enzyme 1) (Beta-site amyloid precursor protein cleaving enzyme 1) (Aspartyl protease 2) (Aap 2) (ASP2) (Membrane-associated aspartic protease 2) (Memapsin-2).

GN Name=BACE1; Synonym=BACE;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 NC NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORM A).
 RC TISSUE=Brain;
 RX MEDLINE=20002972; PubMed=10531052; DOI=10.1126/science.286.5440.735;
 RA Vaasar R., Bennett B.D., Babu-Khan S., Kahn S., Mendiaz E.A.,
 RA Denis P., Teplow D.B., Ross S., Amaranse P., Loeffler R., Luo Y.,
 RA Fisher S., Puller J., Edenson S., Lile J., Jarosinski M.A.,
 RA Biere A.L., Curran E., Burgess T., Louis J.-C., Collins F.,
 RA Treanor J., Rogers G., Citron M.;
 RT "Beta-secretase cleavage of Alzheimer's amyloid precursor protein by
 the transmembrane aspartic protease BACE.";
 RL Science 286:735-741(1999).
 RN [2]
 RP SEQUENCE FROM N.A. (ISOFORM A), SEQUENCE OF 46-68, AND
 RC CHARACTERIZATION.
 RX TISSUE=Brain;
 RX MEDLINE=20057171; PubMed=10591214; DOI=10.1038/990114;
 RA Sinha S., Anderson J.P., Barbour R., Basl G.S., Caccavello R.,
 RA Davis D., Dean M., Doney H.F., Frigon N., Hong J., Jacobson-Croak K.,
 RA Jewett N., Keim P., Knops J., Lieberburg I., Power M., Tan H.,
 RA Tatsuno G., Tung J., Schenk D., Seubert P., Suematsaari S.M., Wang S.,
 RA Walker D., Zhao J., McConlogue L., Varghese J.;
 RT "Purification and cloning of amyloid precursor protein beta-secretase
 from human brain.";
 RL Nature 402:537-540(1999).
 RN [3]
 RP SEQUENCE FROM N.A. (ISOFORM A).
 RX MEDLINE=20057170; PubMed=10591213; DOI=10.1038/990107;
 RA Yan R., Bienkowski M.J., Shuck M.E., Miao H., Torry M.C., Pauley A.M.,
 RA Braahler J.R., Stratan N.C., Mathews W.R., Buhl A.E., Carter D.B.,
 RA Tomasselli A.G., Parodi L.A., Heinrichson R.L., Gurney M.E.;
 RT "Membrane-anchored aspartyl protease with Alzheimer's disease beta-
 secretase activity.";
 RL Nature 402:533-537(1999).
 RN [4]
 RP SEQUENCE FROM N.A. (ISOFORM A).
 RX MEDLINE=20120043; PubMed=10656250; DOI=10.1006/mcne.1999.0811;
 RA Hussain I., Powell D.J., Howlett D.R., Tew D.G., Meek T.D.,
 RA Chapman C., Gloger I.S., Murphy K.E., Southern C.D., Ryan D.M.,
 RA Smith T.S., Simons D.L., Walsh F.S., Dingwall C., Christie G.;
 RT "Identification of a novel aspartic protease (Asp 2) as beta-
 secretase.";
 RL Mol. Cell. Neurosci. 14:419-427(1999).
 RN [5]
 RP SEQUENCE FROM N.A. (ISOFORM B).
 RC TISSUE=Brain, and Pancreas;
 RA Michel B., De Pietri Tonelli D., Zaccchetti D., Keller P.;
 RT "New beta-site APP cleaving enzyme isoform (BACE-1B) obtained from
 human brain and pancreas.";
 RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
 RN [6]
 RP SEQUENCE FROM N.A. (ISOFORM C).
 RC TISSUE=Pancreas;
 RA Zaccchetti D., De Pietri Tonelli D., Schnutbus R.;
 RT "New beta-site APP cleaving enzyme isoform (BACE-1C) obtained from
 human pancreas.";
 RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
 RN [7]
 RP SEQUENCE FROM N.A. (ISOFORMS B; C AND D).
 RC TISSUE=Brain;
 RX MEDLINE=21408467; PubMed=11516562; DOI=10.1016/S0304-3940(01)01912-7;
 RA Tanahashi H., Tsubita T.;
 RT "Three novel alternatively spliced isoforms of the human beta-site
 amyloid precursor protein cleaving enzyme (BACE) and their effect on
 amyloid beta-peptide production.";
 RL Neurosci. Lett. 307:9-12(2001).
 RN [8]
 RP SEQUENCE OF 14-501 FROM N.A. (ISOFORM A), AND CHARACTERIZATION.

RX MEDLINE=20144060; PubMed=10677483; DOI=10.1073/pnas.97.4.1456;
 RA Lin X., Koelsch G., Wu S., Downs D., Dachtel A., Tang J.;
 RT "Human aspartic protease memapsin 2 cleaves the beta-secretase site of
 beta-amyloid precursor protein.";
 RL Proc. Natl. Acad. Sci. U.S.A. 97:1456-1460(2000).
 RN [9]
 RP DISULFIDE BONDS.
 RX MEDLINE=21950860; PubMed=11953458;
 RA Fischer F., Molinari M., Bodendorf U., Paganetti P.;
 RT "The disulphide bonds in the catalytic domain of BACE are critical but
 not essential for amyloid precursor protein processing activity.";
 RL J. Neurochem. 80:1079-1086(2002).
 CC -1- FUNCTION: Responsible for the proteolytic processing of the
 amyloid precursor protein (APP). Cleaves at the amino terminus of
 the A-beta peptide sequence, between residues 671 and 672 of APP,
 leads to the generation and extracellular release of beta-cleaved
 soluble APP, and a corresponding cell-associated carboxy-terminal
 fragment which is later released by gamma-secretase.
 CC -1- CATALYTIC ACTIVITY: Broad endopeptidase specificity. Cleaves Glu-
 Val-Asn-Leu-I-Asp-Ala-Glu-Phe in the Swedish variant of
 Alzheimer's amyloid precursor protein.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 Event=Alternative splicing; Named isoforms=4;
 Name=A; Synonyms=BACE-1A, BAC-501,
 IsoId=P56817-1; Sequence=Displayed;
 Name=B; Synonyms=BACE-1B, BACE-I-476;
 IsoId=P56817-2; Sequence=VSP_005223;
 Name=C; Synonyms=BACE-1C, BACE-I-457;
 IsoId=P56817-3; Sequence=VSP_005222;
 Name=D; Synonyms=BACE-1D, BACE-I-432;
 IsoId=P56817-4; Sequence=VSP_005222; VSP_005223;
 CC -1- TISSUE SPECIFICITY: Brain.
 CC -1- SIMILARITY: Belongs to the peptidase A1 family.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; AF190725; AAF04142.1; -
 DR EMBL; AF201468; AAF18982.1; -
 DR EMBL; AF200343; AAF17079.1; -
 DR EMBL; AF204943; AAF26367.1; -
 DR EMBL; AF338816; AAK38374.1; -
 DR EMBL; AF338817; AAK38375.1; -
 DR EMBL; AB050436; BAB40931.1; -
 DR EMBL; AB050437; BAB40932.1; -
 DR EMBL; AB050438; BAB40933.1; -
 DR EMBL; AF200193; AAF13715.1; -
 DR PIR; A59090; A59090
 DR PDB; 1FKN; X-ray; A/B=56-446.
 DR PDB; 1M4H; X-ray; A/B=56-446.
 DR MEROPS; A01.004; -
 DR Genew; HGNC:933; BACE1.
 DR H-InvDB; HIX0010165; -
 DR MIM; 604252; -
 DR GO; GO:0005887; C:integral to plasma membrane; TAS.
 DR GO; GO:0008798; F:beta-aspartyl-peptidase activity; TAS.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; TAS.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009120; Pept_A1_BACE.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR InterPro; IPR009007; Pept_AspArtic.
 DR InterPro; IPR001461; Peptidase_A1.
 DR Pfam; PF00026; Asp_1.
 DR PRINTS; PRO1816; BACE1.
 DR PRINTS; PRO1815; BACEFAMILY.
 DR PRINTS; PRO0792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.

KW 3D-structure; Alternative splicing; Aspartyl protease;
 KW Direct protein sequencing; Glycoprotein; Hydrolase; Signal;
 KW Transmembrane; Zymogen.
 FT SIGNAL 1 21 Potential.
 FT PROPEP 22 45 Beta-secretase 1.
 FT CHAIN 46 501 Extracellular (Potential).
 FT DOMAIN 22 457 Potential.
 FT TRANSMEM 458 478 Cytoplasmic (Potential).
 FT DOMAIN 479 501 By similarity.
 FT ACT_SITE 93 93 By similarity.
 FT ACT_SITE 289 289 By similarity.
 FT DISULFID 216 420
 FT DISULFID 278 443
 FT DISULFID 330 380
 FT CARBOHYD 153 153 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 172 172 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 223 223 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 354 354 N-linked (GlcNAc...) (Potential).
 FT VARSPLIC 146 189 Missing (in isoform C and isoform D).
 FT VARSPLIC 190 214 /FTid=VSP_005222.
 FT VARSPLIC 214 214 Missing (in isoform B and isoform D).
 FT VARSPLIC 214 214 /FTid=VSP_005223.
 FT HELIX 61 63
 FT TURN 64 65
 FT STRAND 67 70
 FT TURN 71 73
 FT STRAND 74 81
 FT TURN 82 85
 FT STRAND 86 93
 FT TURN 94 95
 FT STRAND 99 102
 FT TURN 107 108
 FT HELIX 115 117
 FT TURN 119 120
 FT STRAND 122 131
 FT STRAND 136 147
 FT TURN 149 150
 FT STRAND 155 167
 FT TURN 172 173
 FT STRAND 178 181
 FT HELIX 185 187
 FT TURN 192 193
 FT HELIX 197 204
 FT STRAND 211 215
 FT HELIX 224 229

Query Match 100.0%; Score 16; DB 1; Length 501;
 Best Local Similarity 100.0%; Pred. No. 4.2e-08;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TOHGIRLPURSGIGA 16
 DB 22 TOHGIRLPURSGIGA 37

RESULT 3
 ID Q81YC8 PRELIMINARY; PRT; 501 AA.
 AC Q81YC8;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Beta-site APP-cleaving enzyme 1, isoform A preproprotein.
 GN Name=BACE1;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OC NCB1_TaxID=9606;
 OX NCB1_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Strauberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,

RA Klausner R.D., Collins F.S., Wagner L., Sherman C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carrinci P., Prange C.,
 RA Raha S.S., Loguella N.A., Peters G.J., Abramson R.D., Mullany S.J.,
 RA Bosak S.A., McGowan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Morley K.C., Hale S., Garcia A.M., Gay L.J., Huiyk S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kerteman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butlerfield Y.S.,
 RA Krzywinski M.I., Skalska U., Smailus D.E., Scherch A., Schein J.E.,
 RA Jones S.J., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RA Strauberg R.;
 RL Submitted (JUL-2002) to the EMBL/Genbank/DBJ databases.
 CC -1 SIMILARITY: Belongs to peptidase family A1.
 DR EMBL; BC036084; AAH36084.1; -.
 DR HSP; P56817; 1PKN.
 DR GO; GO:0005768; C:cytosol; ISS.
 DR GO; GO:0005794; C:Golgi apparatus; ISS.
 DR GO; GO:0016021; C:integral to membrane; ISS.
 DR GO; GO:0004190; F:aspartic-type endopeptidase activity; ISS.
 DR GO; GO:0050435; P:beta-amyloid metabolism; ISS.
 DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro; IPR001461; Peptidase A1.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009120; Pept_A1_BACE1.
 DR InterPro; IPR009007; Pept_Aspartic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PRINTS; PRO1816; BACE1.
 DR PRINTS; PRO1815; BACEFAMILY.
 DR PRINTS; PRO0792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.
 DR Aspartyl protease; Hydrolase; Protease.
 SQ SEQUENCE 501 AA; 55823 MW; 768595CF5517EB7 CRC64;

Query Match 100.0%; Score 16; DB 2; Length 501;
 Best Local Similarity 100.0%; Pred. No. 4.2e-08;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TOHGIRLPURSGIGA 16
 DB 22 TOHGIRLPURSGIGA 37

RESULT 4
 ID Q8CAF4 PRELIMINARY; PRT; 467 AA.
 AC Q8CAF4;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Mus musculus 0 day neonate cerebellum cDNA, RIKEN full-length enriched
 DE library, clone: C230037B16 product: beta-site APP cleaving enzyme, full
 DE insert sequence.
 GN Name=Bace1; Synonym=Bace;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OC NCB1_TaxID=10090;
 OX NCB1_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=99279253; PubMed=10349636; DOI=10.1016/S0076-6879(99)03004-9;
 RA

RA Carninci P., Hayashizaki Y.;
 RT "High-efficiency full-length cDNA cloning";
 RL Meth. Enzymol. 303:19-44(1999).
 RN (2)
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=21085660; PubMed=11217851; DOI=10.1038/35055500;
 RA RIKEN FANTOM Consortium;
 RT "Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 409:685-690(2001).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RA The FANTOM Consortium;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RL Nature 420:563-573(2002).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=20499374; PubMed=11042159; DOI=10.1101/gr.145100;
 RA Carninci P., Shibata Y., Hayatsu N., Sugahara Y., Shibata K., Itoh M.,
 RT Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
 RL "Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new genes.";
 RL Genome Res. 10:1617-1630(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=20530913; PubMed=11076861; DOI=10.1101/gr.152600;
 RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
 RT Kono H., Akiyama J., Nishi K., Kikunishi T., Tashiro H., Itoh M.,
 RA Sumi N., Ishii Y., Nakamura S., Hazama M., Nishino T., Harada A.,
 RA Yamamoto R., Matsumoto H., Sakaguchi S., Ikegami T., Kashiwagi K.,
 RA Fujiwaka S., Inoue K., Togawa K., Izawa M., Ohara E., Watanabe M.,
 RA Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsura S., Kawai J.,
 RA Okazaki Y., Muramatsu M., Inoue Y., Kira A., Hayashizaki Y.;
 RT "RIKEN integrated sequence analysis (RISA) system-384-format
 RL sequencing pipeline with 384 multicapillary sequencer.";
 RL Genome Res. 10:11757-1771(2000).
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RA Adachi J., Aizawa K., Akiyama T., Aizawa T., Bono H., Carninci P.,
 RA Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
 RA Hayashida K., Hayatsu N., Hiramoto K., Hiraoka T., Hirozane T.,
 RA Hori F., Imotani K., Ishii Y., Itoh M., Kagawa I., Kasukawa T.,
 RA Katoh H., Kawai J., Kojima Y., Kondo S., Kono H., Kouda M., Koya S.,
 RA Kurihara C., Matsuyama T., Miyazaki R., Murata M., Nakamura M.,
 RA Nishii K., Nomura K., Numazaki R., Ohno M., Ohsato N., Okazaki Y.,
 RA Saito R., Saitoh H., Sakai C., Sakai K., Sakazume N., Sano H.,
 RA Sasaki D., Shibata K., Shinagawa A., Shiraki T., Sogabe Y., Tagami M.,
 RA Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
 RA Tomaru A., Toya T., Yasunishi A., Muramatsu M., Hayashizaki Y.;
 RL Submitted (Apr-2002) to the EMBL/Genbank/DBJ databases.
 CC -1 SIMILARITY: Belongs to peptidase family A1.
 DR EMBL; AK082317; BAC38462.1; -
 DR HSSP; P56817; IPKN.
 DR MGD; MGI:1346542; Bace1.
 DR GO; GO:0005768; C:cytosol; ISS.
 DR GO; GO:0005615; C:extracellular space; TAS.
 DR GO; GO:0005794; C:Golgi apparatus; ISS.
 DR GO; GO:0016021; C:integral to membrane; ISS.
 DR GO; GO:0004190; F:aspartic-type endopeptidase activity; ISS.
 DR GO; GO:0050435; F:beta-amyloid metabolism; ISS.
 DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro; IPR001461; Peptidase A1.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009120; Pept_A1_BACE1.
 DR InterPro; IPR009007; Pept_Aspartic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PRINTS; PRO1816; BACE1.

DR PRINTS; PRO1815; BACEFAMILY.
 DR PRINTS; PRO0792; PEPSTATIN.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.
 KW Aspartyl protease; Hydrolase; Protease.
 SQ SEQUENCE 467 AA; 52063 MW; 31AB674FF1843652 CRC64;
 Query Match 62.5%; Score 10; DB 2; Length 467;
 Best local similarity 100.0%; Pred. No. 0.05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 4 GIRLPRLRSL 13
 DB 25 GIRLPRLRSL 34
 AC P56818;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 25-OCT-2004 (Rel. 45, Last annotation update)
 DE Beta-secretase 1 precursor (EC 3.4.23.46) (Beta-site APP cleaving
 DE enzyme 1) (Beta-site amyloid precursor protein cleaving enzyme 1)
 DE (Aspartyl protease 2) (Asp 2) (ASP2) (Membrane-associated aspartic
 DE protease 2) (Memapsin-2).
 GN Name=BACE1; Synonyms=Bace;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20002972; PubMed=10531052; DOI=10.1126/science.286.5440.735;
 RA Vassar R., Bennett B.D., Babu-Khan S., Kahn S., Mendez E.A.,
 RA Denis P., Teplow D.B., Ross S., Amarante P., Loefler R., Luo Y.,
 RA Fisher S., Fuller J., Edenson S., Lile J., Jarosinski M.A.,
 RA Biere A.L., Curran E., Burgess T., Louis J.-C., Collins F.,
 RA Treanor D., Rogers G., Citron M.;
 RT "Beta-secretase cleavage of Alzheimer's amyloid precursor protein by
 RT the transmembrane aspartic protease BACE.";
 RL Science 286:735-741(1999).
 RN [2]
 RP SEQUENCES TO 6 AND 81-87.
 RA Bennett B.D., Vassar R., Citron M.;
 RL Submitted (Jan-2000) to the EMBL/Genbank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20057170; PubMed=10591213; DOI=10.1038/990107;
 RA Yan R., Bienkowski M.J., Shuck M.E., Miao H., Toriy M.C., Paulay A.M.,
 RA Braehler J.R., Strattan N.C., Mathews W.R., Buhl A.E., Carter D.B.,
 RA Tomasselli A.G., Parodi L.A., Heinrichson R.L., Gurney M.E.;
 RT "Membrane-associated aspartyl protease with Alzheimer's disease beta-
 RT secretase activity.";
 RL Nature 402:533-537(1999).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Head;
 RX MEDLINE=22354683; PubMed=12466851; DOI=10.1038/nature01266;
 RA Okazaki Y., Furuno M., Kasukawa T., Adachi J., Bono H., Kondo S.,
 RA Nishida I., Oseko N., Saito R., Suzuki H., Yamanaka I., Kiyosawa H.,
 RA Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Schonbach G., Gotohori T.,
 RA Badarrelli R., Hill D.P., Bull C., Hume D.A., Quackenbush J.,
 RA Schmitt L.M., Kanapin A., Matsuda H., Batelov S., Beisel K.W.,
 RA Blake J.A., Bradt D., Brusic V., Chochia C., Corbani L.E., Cousins S.,
 RA Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Fraser K.S.,
 RA Gaasterland T., Gariboldi M., Gissi C., Godzik A., Gough J.,
 RA Grimmerond S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D.,
 RA Kanai A., Kawai H., Kawasawa Y., Kedzierski R.M., King B.L.,
 RA Kanagaya A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A.,
 RA Maglott D.R., Malais L., Marchionni L., McKenzie L., Mikki H.,
 RA Nagashima T., Nomura K., Okada T., Pavan W.J., Petrea G., Pesole G.,
 RA Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,

RA Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
RA Sandelin A., Schneider C., Sempé C.A., Setou M., Shindada K.,
RA Sultana R., Takenaka Y., Taylor M.S., Teasdale R.D., Tomita M.,
RA Verardo R., Wagner L., Wahlestedt C., Wang Y., Watanabe Y., Wells C.,
RA Wilm L.G., Wymshaw-Boris A., Yanagisawa M., Yang L., Yang L.,
RA Yuan Z., Zevonci M., Zhu Y., Zimmer A., Carninci P., Hayatsu N.,
RA Hirozane-Kishikawa T., Kono H., Nakamura M., Sakazume N., Sato K.,
RA Shitaki T., Naki K., Kawai J., Aizawa K., Arakawa T., Fukuda S.,
RA Hara A., Hashizume W., Imotani K., Ishii Y., Itoh M., Kagawa I.,
RA Miyazaki A., Sakai K., Sasaki D., Shibata K., Shingaga A.,
RA Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J.,
RA Birney E., Hayashizaki Y.,
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs";
RN Nature 420:563-573(2002).
RP [5]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6; TISSUE=Brain;
RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Scheefter C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stoplecko M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Ueda T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mallary S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulik S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kerteman M., Madan A.C., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A.C., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butcherfield Y.S.N., Krzywinski M.I., Skalska U., Smalins D.E.,
RA Scherch A., Schein J.E., Jones S.J.M., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -1- FUNCTION: Responsible for the proteolytic processing of the
CC amyloid precursor protein (APP). Cleaves at the amino terminus of
CC the A-beta peptide sequence, between residues 671 and 672 of APP,
CC leads to the generation and extracellular release of beta-cleaved
CC soluble APP, and a corresponding cell-associated carboxy-terminal
CC fragment which is later released by gamma-secretase (By
CC similarity).
CC -1- CATALYTIC ACTIVITY: Broad endopeptidase specificity. Cleaves Glu-
CC Val-Asn-Ileu-[Asp-Ala-Glu-Phe in the Swedish variant of
CC Alzheimer's amyloid precursor protein.
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- TISSUE SPECIFICITY: Brain.
CC -1- SIMILARITY: Belongs to the peptidase A1 family.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AF190726; AAF04143.2; -;
CC EMBL: AF200346; AAF17082.1; -;
CC EMBL: AK014464; BAB29370.1; -;
CC EMBL: BC048189; AAH48189.1; -;
CC HSSP: P56817; 1M4H.
CC MEROPS: A01.004; -;
CC MGD: MGI:1346542; Bace1.
CC InterPro: IPR009119; Pept_A1_BACE.
CC InterPro: IPR009120; Pept_A1_BACE1.
CC InterPro: IPR001969; Pept_A1_AS.
CC InterPro: IPR009007; Pept_Aspartic.
CC InterPro: IPR001461; Peptidase_A1.

DR Pfam: PF00026; Asp. 1.
DR PRINTS: PRO1816; BACE1.
DR PRINTS: PRO1815; BACEFAMILY.
DR PRINTS: PRO0792; PEPsin.
DR PROSITE: PS00141; ASP-PROTEASE, 1.
KW Aspartyl protease; Glycoprotein; Hydrolase; Signal; Transmembrane;
KW Zymogen.
FT SIGNAL 1 21 Potential.
FT PROPEP 22 45 Potential.
FT CHAIN 46 501 Beta-secretase 1.
FT DOMAIN 22 457 Extracellular (Potential).
FT TRANSMEM 458 478 Cytoplasmic (Potential).
FT DOMAIN 479 501 Potential.
FT ACT_SITE 93 93 By similarity.
FT ACT_SITE 289 289 By similarity.
FT DISULFID 216 420 By similarity.
FT DISULFID 278 443 By similarity.
FT DISULFID 330 380 By similarity.
FT CARBOHYD 153 153 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 172 172 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 223 223 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 354 354 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 501 AA; 55747 MW; C085A013145E474E CRC64;
Query Match 62.5%; Score 10; DB 1; Length 501;
Best Local Similarity 100.0%; Pred. No. 0.053;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 4 GIRLPRLSGL 13
Db 25 GIRLPRLSGL 34
RESULT 6
BAEL RAT STANDARD; PRT; 501 AA.
ID BAEL RAT
AC P56819;
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 30-MAY-2000 (Rel. 39, Last annotation update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE beta-secretase 1 precursor (EC 3.4.23.46) (Beta-site APP cleaving
DE enzyme 1) (Beta-site amyloid precursor protein cleaving enzyme 1)
DE (Aspartyl) protease 2) (Asp 2) (ASP2) (Membrane-associated aspartic
DE protease 2) (Memapsin-2).
GN Name=Bace1; Synonyms=Bace;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20002972; PubMed=10531052; DOI=10.1126/science.286.5440.735;
RA Vassar R., Bennett B.D., Babu-Khan S., Kahn S., Mendiaz E.A.,
RA Denis P., Teplow D.B., Ross S., Amarante P., Loefler R., Luo Y.,
RA Fisher S., Fuller J., Edenson S., Lile J., Jarosinski M.A.,
RA Biere A.L., Curran E., Burgess T., Louis J.-C., Collins F.,
RA Treanor J., Rogers G., Citron M.,
RT "Beta-secretase cleavage of Alzheimer's amyloid precursor protein by
RT the transmembrane aspartic protease BACE";
RL Science 286:735-741(1999).
CC -1- FUNCTION: Responsible for the proteolytic processing of the
CC amyloid precursor protein (APP). Cleaves at the amino terminus of
CC the A-beta peptide sequence, between residues 671 and 672 of APP,
CC leads to the generation and extracellular release of beta-cleaved
CC soluble APP, and a corresponding cell-associated carboxy-terminal
CC fragment which is later released by gamma-secretase (By
CC similarity).
CC -1- CATALYTIC ACTIVITY: Broad endopeptidase specificity. Cleaves Glu-
CC Val-Asn-Ileu-[Asp-Ala-Glu-Phe in the Swedish variant of
CC Alzheimer's amyloid precursor protein.
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- SIMILARITY: Belongs to the peptidase A1 family.
CC -----

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 CC or send an email to license@isb-eb.ch).

DR EMBL: AF190727; AF04144.1; -
 DR HSSP: P56817; IM4H.
 DR MEROPS: A01.004; -
 DR RGD: 2191; Bace.
 DR InterPro: IPR009119; Pept_A1_BACE.
 DR InterPro: IPR009120; Pept_A1_BACE1.
 DR InterPro: IPR009169; Pept_Asp_AS.
 DR InterPro: IPR009007; Pept_AspArtic.
 DR InterPro: IPR001461; Peptidase_A1.
 DR Pfam: PF00026; Asp.1.
 DR PRINTS: PRO1816; BACE1.
 DR PRINTS: PRO1815; BACEFAMILY.
 DR PRINTS: PRO0792; PEPSTN.
 DR PROSITE: PS00141; ASP_PROTEASE; 1.
 DR Aspartyl protease; Glycoprotein; Hydrolase; Signal; Transmembrane;
 KM zymogen.
 FT SIGCAT: 1 21
 FT PROPEP: 22 45
 FT CHAIN: 46 501
 FT DOMAIN: 22 457
 FT TRANSMEM: 458 478
 FT DOMAIN: 479 501
 FT ACT SITE: 93 93
 FT ACT SITE: 289 289
 FT DISULFID: 216 420
 FT DISULFID: 278 443
 FT DISULFID: 310 380
 FT CARBOHD: 153 153
 FT CARBOHD: 172 172
 FT CARBOHD: 223 223
 FT CARBOHD: 354 354
 FT CARBOHD: 501 AA; 55806 MW; 248445BC8B87DE3 CRC64;
 SQ SEQUENCE

Query Match
 Best Local Similarity 62.5%; Score 10; DB 1; Length 501;
 Matches 10; Conservativity 100.0%; Pred. No. 0.053; Mismatches 0; Indels 0; Gaps 0;

Qy 4 GIRLPLRSGL 13
 Db 25 GIRLPLRSGL 34

RESULT 7
 O8BOY4 PRELIMINARY; PRT; 501 AA.
 AC O8BOY4; 01-MAR-2003 (Tremblrel. 23. Created)
 DT 01-MAR-2003 (Tremblrel. 23. Last sequence update)
 DE Mus musculus adult male corpora quadrigemina update)
 DE enriched library; clone: B230346M13 product: beta-site APP cleaving
 DE enzyme; full insert sequence.
 GN Name=Bace1; Synonyms=Bace;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=99279253; PubMed=10349636; DOI=10.1016/S0076-6879(99)03004-9;
 RA Carninci P.; Hayashizaki Y.;
 RT "High-efficiency full-length cDNA cloning";
 RU Meth. Enzymol. 303:19-44(1999).
 RN [2]

RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=21085660; PubMed=11217851; DOI=10.1038/35055500;
 RA RIKEN FANTOM Consortium;
 RT "Functional annotation of a full-length mouse cDNA collection";
 RU Nature 409:685-690(2001).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RA The FANTOM Consortium;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs";
 RU Nature 420:563-573(2002).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=20499374; PubMed=11042159; DOI=10.1101/gr.145100;
 RA Carninci P.; Shibata Y.; Hayatsu N.; Sugahara Y.; Shibata K.; Itoh M.;
 RT "Normalization and subcloning of cap-trapper-selected cDNAs to
 RT prepare full-length cDNA libraries for rapid discovery of new genes";
 RU Genome Res. 10:1617-1630(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=20530913; PubMed=11076861; DOI=10.1101/gr.152600;
 RA Shibata K.; Itoh M.; Aizawa K.; Nagaoka S.; Sasaki N.; Carninci P.;
 RA Kono H.; Akiyama U.; Nishi K.; Kitsuana T.; Tashiro H.; Itoh M.;
 RA Sumi N.; Ishii Y.; Nakamura S.; Hazama M.; Nishine T.; Harada A.;
 RA Yamamoto R.; Matsumoto H.; Sakaguchi S.; Ikegami T.; Kashiwagi K.;
 RA Fujiwaka S.; Inoue K.; Togawa Y.; Izawa M.; Ohara E.; Wataniki M.;
 RA Yoneda Y.; Ishikawa T.; Ozawa K.; Tanaka T.; Matsushita S.; Kawai J.;
 RA Okazaki Y.; Muramatsu M.; Inoue Y.; Kira A.; Hayashizaki Y.;
 RT RIKEN integrated sequence analysis (RISA) system-384-format
 RT sequencing pipeline with 384 multicapillary sequencer";
 RU Genome Res. 10:1757-1771(2000).
 RN [6]

RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RA Adachi U.; Aizawa K.; Akimura T.; Arakawa T.; Bono H.; Carninci P.;
 RA Fukuda S.; Furuno M.; Hanagaki T.; Hara A.; Hashizume W.;
 RA Hayashida K.; Hayatsu N.; Hiramoto K.; Hiraoka T.; Hirozane T.;
 RA Hori F.; Imocani K.; Ishii Y.; Itoh M.; Kaga S.; Kondo S.; Kouda M.; Koyama S.;
 RA Kurihara C.; Matsuyama T.; Miyazaki R.; Murata M.; Nakamura M.;
 RA Nihi K.; Nomura K.; Numazaki R.; Ohno M.; Ohgaki N.; Okazaki Y.;
 RA Saito R.; Saitoh H.; Sakai C.; Sakai K.; Sakazume N.; Sano H.;
 RA Tagawa A.; Takahashi F.; Shinagawa A.; Shiraki T.; Sogabe Y.; Tagami M.;
 RA Tomaru A.; Toya T.; Yasunishi A.; Muramatsu M.; Hayashizaki Y.;
 RL Submitted (JUL-2001) to the EMBL/Genbank/DBJ databases.
 CC -1- SIMILARITY: Belongs to peptidase family A1.
 DR EMBL: AK046175; BAC32620.1; -
 DR HSSP: P56817; 1FKV
 DR MGD: MGI:1346542; Bace1.
 DR GO: GO:0005768; C:cytosol; ISS.
 DR GO: GO:0005615; C:extracellular space; TAS.
 DR GO: GO:0005794; C:Golgi apparatus; ISS.
 DR GO: GO:0016021; C:integral to membrane; ISS.
 DR GO: GO:0004190; F:aspartic-type endopeptidase activity; ISS.
 DR GO: GO:0050435; F:beta-amyloid metabolism; ISS.
 DR GO: GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro: IPR001461; Peptidase_A1.
 DR InterPro: IPR009119; Pept_A1_BACE.
 DR InterPro: IPR009120; Pept_A1_BACE1.
 DR InterPro: IPR009007; Pept_AspArtic.
 DR InterPro: IPR001969; Pept_Asp_AS.
 DR PRINTS: PRO1816; BACE1.
 DR PRINTS: PRO1815; BACEFAMILY.
 DR PRINTS: PRO0792; PEPSTN.
 DR PROSITE: PS00141; ASP_PROTEASE; 1.
 KM Aspartyl protease; Hydrolase; Protease.

SQ SEQUENCE 501 AA; 55816 MW; C0855513145E024E CRC64;
 Query Match 62.5%; Score 10; DB 2; Length 501;
 Best Local Similarity 100.0%; Pred. NO. 0.053;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GIRLPLRSGL 13
 |||||
 DB 25 GIRLPLRSGL 34

RESULT 8
 Q8C7R1 PRELIMINARY; PRT; 501 AA.
 ID Q8C7R1
 AC Q8C7R1; 01-MAR-2003 (TREMBlrel. 23, Created)
 DT 01-MAR-2003 (TREMBlrel. 23, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Mus musculus 12 days embryo spinal cord cDNA, RIKEN full-length
 DE enriched library, clone: C530008K17 product: beta-site APP cleaving
 DE enzyme, full insert sequence.
 GN Name=Bacel; Synonyms=Bace;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OC NCBI_TaxID=10090;
 [1]
 RN SEQUENCE FROM N.A.
 RP STRAIN=C57BL/6J; TISSUE=spinal cord;
 RC MEDLINE=99279253; PubMed=10349636; DOI=10.1016/S0076-6879(99)03004-9;
 RX RIKEN FANTOM Consortium;
 RA "Functional annotation of a full-length mouse cDNA collection."
 RT Nature 409:685-690 (2001).
 RL [3]
 RN SEQUENCE FROM N.A.
 RP STRAIN=C57BL/6J; TISSUE=spinal cord;
 RC The FANTOM Consortium;
 RA the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs."
 RL Nature 420:563-573 (2002).
 [4]
 RN SEQUENCE FROM N.A.
 RP STRAIN=C57BL/6J; TISSUE=spinal cord;
 RC MEDLINE=20499374; PubMed=11042159; DOI=10.1101/gr.145100;
 RX Karimci P., Shibata Y., Hayatsu M., Sugahara Y., Shibata K., Itoh M.,
 RA Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
 RT "Normalization and substructure of cap-trapper-selected cDNAs to
 RT prepare full-length cDNA libraries for rapid discovery of new genes."
 RL Genome Res. 10:1617-1630 (2000).
 [5]
 RN SEQUENCE FROM N.A.
 RP STRAIN=C57BL/6J; TISSUE=spinal cord;
 RC MEDLINE=20530913; PubMed=11076861; DOI=10.1101/gr.152600;
 RX Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
 RA Kono H., Akiyama J., Nishi K., Katsunai T., Tashiro H., Itoh A.,
 RA Sumi N., Ishii Y., Nakamura S., Hazama M., Nishino T., Harada A.,
 RA Yamamoto R., Matsumoto H., Sakaguchi S., Ikegami T., Kashiyagi K.,
 RA Fujisake S., Inoue K., Togawa Y., Izawa M., Ohara E., Watabiki M.,
 RA Toneya Y., Ishikawa T., Ozawa K., Tanaka T., Matsuda S., Kawai J.,
 RA Okazaki Y., Muramatsu M., Inoue Y., Kita A., Hayashizaki Y.;
 RT RIKEN integrated sequence analysis (RISA) system-384-format
 RT sequencing pipeline with 384 multicapillary sequencer."
 RL Genome Res. 10:1757-1771 (2000).
 [6]
 RN SEQUENCE FROM N.A.

RC STRAIN=C57BL/6J; TISSUE=spinal cord;
 RA Adachi J., Aizawa K., Akimura T., Aikawa T., Bono H., Carninci P.,
 RA Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
 RA Hayashida K., Hayatsu N., Hiramoto K., Hiraoaka T., Hirozawa T.,
 RA Horii F., Imotani K., Ishii Y., Itoh M., Kagawa I., Kaikawa T.,
 RA Kato H., Kawai J., Kojima Y., Kondo S., Kono H., Konda M., Koya S.,
 RA Kurihara C., Matsuyama T., Miyazaki A., Murata M., Nakamura M.,
 RA Nishi K., Nomura K., Numazaki R., Ohno M., Ohsato N., Okazaki Y.,
 RA Saito R., Saitoh H., Sakai C., Sakai K., Sakazume N., Sano H.,
 RA Sasaki D., Shibata K., Shinagawa A., Shiraki T., Sogabe Y., Tagami M.,
 RA Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
 RA Tomaru A., Toya T., Yasunishi A., Muramatsu M., Hayashizaki Y.;
 RL Submitted (Jul-2001) to the EMBL/GenBank/DBJ databases.
 CC -1-SIMILARITY: belongs to peptidase family A1.
 DR EMBL; AK049626; BAC33844.1; --
 DR HSSP; P56817; 1FKN.
 DR MGD; MGI:1346542; Bacel.
 DR GO; GO:0005768; C:cytosol; ISS.
 DR GO; GO:0005615; C:extracellular space; TAS.
 DR GO; GO:0005794; C:Golgi apparatus; ISS.
 DR GO; GO:0016021; C:integral to membrane; ISS.
 DR GO; GO:0004190; F:aspartic-type endopeptidase activity; ISS.
 DR GO; GO:0050435; F:beta-amyloid metabolism; ISS.
 DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro: IPR001461; Peptidase A1.
 DR InterPro: IPR009119; Pept_A1_BACE1.
 DR InterPro: IPR009120; Pept_A1_BACE1.
 DR InterPro: IPR009007; Pept_Aspartic.
 DR InterPro: IPR001969; Pept_Asp_AS.
 DR PRINTS; PR01815; BACEFAMILY.
 DR PRINTS; PR00792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.
 DR Aspartyl protease, Hydrolyase; Protease.
 KW SEQUENCE 501 AA; 55761 MW; B410DAB6647663 CRC64;

Query Match 62.5%; Score 10; DB 2; Length 501;
 Best Local Similarity 100.0%; Pred. NO. 0.053;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 GIRLPLRSGL 13
 |||||
 DB 25 GIRLPLRSGL 34

RESULT 9
 Q7S2I8 PRELIMINARY; PRT; 814 AA.
 ID Q7S2I8
 AC Q7S2I8; 01-OCT-2003 (TREMBlrel. 25, Created)
 DT 01-OCT-2003 (TREMBlrel. 25, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE SMC1 alpha (Fragment).
 OS Oryzias latipes (Medaka fish) (Japanese ricefish).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 OC Acanthopterygii; Acanthopterygii; Perciformes; Atherinomorpha;
 OC OC Belontiiformes; Adrianichthyidae; Oryziatidae; Oryzias.
 OC NCBI_TaxID=8090;
 [1]
 RN SEQUENCE FROM N.A.
 RP TISSUE=Testis;
 RC MEDLINE=2260315; PubMed=12759374;
 RX Lee J., Iwai T., Yokota T., Yamashita M.;
 RT "Temporally and spatially selective loss of Rec8 protein from meiotic
 RT chromosomes during mammalian meiosis."
 RL J. Cell Sci. 116:2781-2790 (2003).
 CC -1-SIMILARITY: belongs to the ABC transporter family.
 DR EMBL; AB097255; BAC76893.1; --
 DR HSSP; Q9X0R4; IE69.
 DR GO; GO:0016020; C:membrane; IEA.
 DR GO; GO:0005524; F:ATP binding; IEA.
 DR GO; GO:0042626; F:ATPase activity, coupled to transmembrane m. . .; IEA.

```

DR GO:0007059; P:chromosome segregation; IEA.
DR GO:0006810; P:transport; IEA.
DR InterPro; IPR003439; ABC transporter.
DR InterPro; IPR003405; SMC_C.
DR InterPro; IPR010935; SMC_hinge.
DR Pfam; PF02483; SMC_C; 1.
DR Pfam; PF06470; SMC_hinge; 1.
DR Prodom; PD000006; ABC transporter; 1.
DR Prosite; PS00211; ABC_TRANSPORTER_1; UNKNOWN_1.
KM ATP-binding.
FT NON_TER
SQ SEQUENCE 814 AA; 94132 MW; 8653EC762EC6A5A CRC64;

Query Match
Best Local Similarity 50.0%; Score 8; DB 2; Length 814;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 IRLPLRSG 12
Db 521 IRLPLRSG 528

RESULT 10
Q9JPF6 PRELIMINARY; PRT; 97 AA.
ID Q9JPF6
AC Q9JPF6
DT 01-OCT-2000 (TREMblrel. 15, Created)
DT 01-OCT-2000 (TREMblrel. 15, Last sequence update)
DT 01-OCT-2003 (TREMblrel. 25, Last annotation update)
DE Hypothetical protein rch9.
GN Name=rch9;
OS Neisseria meningitidis.
OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
OC Neisseriaceae; Neisseria.
OX NCBI_TaxID=487;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Z2491;
RX DOI=10.1128/JAI.68.4.2082-2095.2000;
RA Klee S.R., Nassif X., Kusecek B., Werker P., Barrett J.L., Achtman M.,
RA Tinsley C.R.;
RT "Molecular and biological analysis of eight genetic islands that
RT distinguish Neisseria meningitidis from the closely related pathogen
RT Neisseria gonorrhoeae".
RL Infect. Immun. 68:2082-2095(2000).
DR EMBL; AJ391256; CAB71967.1; -.
KM Hypothetical protein.
SQ SEQUENCE 97 AA; 10716 MW; 7EDF863F7B6531F9 CRC64;

Query Match
Best Local Similarity 43.8%; Score 7; DB 2; Length 97;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 LRSGLGG 15
Db 81 LRSGLGG 87

RESULT 11
Q9JTC5 PRELIMINARY; PRT; 125 AA.
ID Q9JTC5
AC Q9JTC5
DT 01-OCT-2000 (TREMblrel. 15, Created)
DT 01-OCT-2000 (TREMblrel. 15, Last sequence update)
DT 01-JUN-2003 (TREMblrel. 24, Last annotation update)
DE Hypothetical protein NMA1874.
GN OrderedlocusNames=NMA1874;
OS Neisseria meningitidis (serogroup A).
OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
OC Neisseriaceae; Neisseria.
OX NCBI_TaxID=65699;
RN [1]

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RP SEQUENCE FROM N.A.
RC STRAIN=Z2491 / Serogroup A / Serotype 4A;
RX MEDLINE=20222556; PubMed=10761919; DOI=10.1038/35006655;
RA Parthali J., Achtman M., James K.D., Bentley S.D., Churcher C.M.,
RA Klee S.R., Kerec J.G., Basham D., Brown D., Chillingworth T.,
RA Davies R.M., Davis P., Devlin K., Felwell T., Hamlin N., Holroyd S.,
RA Jørgensen K., Leather S., Mould S., Mungall K.L., Quail M.A.,
RA Rajandream M.A., Rutherford K.M., Simmonds M., Skelton J.,
RA Whitehead S., Spratt B.G., Barrett B.G.;
RT "Complete DNA sequence of a serogroup A strain of Neisseria
RT meningitidis Z2491".
RL Nature 404:502-506(2000).
DR EMBL; AL162757; CAB85097.1; -.
DR PIR; E81814; E81814.
KM Complete proteome; Hypothetical protein.
SQ SEQUENCE 125 AA; 13950 MW; 5B9C7D782E89884D CRC64;

Query Match
Best Local Similarity 43.8%; Score 7; DB 2; Length 125;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 LRSGLGG 15
Db 109 LRSGLGG 115

RESULT 12
Q7NTK4 PRELIMINARY; PRT; 181 AA.
ID Q7NTK4
AC Q7NTK4
DT 01-MAR-2004 (TREMblrel. 26, Created)
DT 01-MAR-2004 (TREMblrel. 26, Last sequence update)
DT 01-MAR-2004 (TREMblrel. 26, Last annotation update)
DE Cytochrome b561.
GN Name=cybB; OrderedlocusNames=CV3050;
OS Chromobacterium violaceum.
OC Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
OC Neisseriaceae; Chromobacterium.
OX NCBI_TaxID=536;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 12472 / DSM 30191;
RX MEDLINE=22882880; PubMed=14500782; DOI=10.1073/pnas.1832124100;
RA Vasconcelos A.T.R., de Almeida D.F., Hungria M., Guimarães C.T.,
RA Antonio R.V., Almeida F.C., de Almeida L.G.P., de Almeida R.,
RA Alves-Gomes J.A., Andrade E.M., Arrupe J., de Araujo M.F.F.,
RA Astolfi-Filho S., Azevedo V., Baptista A.J., Batista L.A.M.,
RA Batista U.S., Belo A., van den Berg C., Bogo M., Bonatto S.,
RA Bordignon J., Brígido M.M., Brito C.A., Brocchi M., Burtly H.A.,
RA Camargo A.A., Cardoso D.D.P., Carneiro N.P., Carraro D.M.,
RA Carvalho C.M.B., Cascardo J.C.M., Cavada B.S., Chuelre L.M.O.,
RA Creczynski-Pasa T.B., Cunha-Junior N.C., Fagundes N., Falcao C.L.,
RA Fantiucci F., Fairies I.P., Felipe M.S.S., Ferrari L.P., Ferro J.A.,
RA Ferro M.I.T., Franco G.R., Freitas N.S.A., Furian L.R.,
RA Gazzinelli R.T., Gomes E.A., Gonçalves P.R., Grangelito T.B.,
RA Gracatapaglia D., Gristard E.C., Hanna E.S., Jardim S.N., Laurino J.,
RA Leoi L.C.T., Lima L.F.A., Loureiro M.F., Lyra M.C.C.P.,
RA Madeira H.M.F., Manfio G.P., Maranhão A.O., Martins W.S.,
RA di Mauro S.M.Z., de Medeiros S.R.B., Meisner R.V., Moreira M.A.M.,
RA Nascimento F.F., Nicolas M.F., Oliveira J.G., Oliveira S.,
RA Paixão R.F.C., Parente J.A., Pedrosa F.O., Pena S.D.J., Pereira J.O.,
RA Pereira M., Pinto L.S.R.C., Pinto L.S., Porto J.I.R., Potrich D.P.,
RA Ramalho-Neto C.E., Reis A.M.M., Rigo L.U., Rondinelli E.,
RA Santos E.B.P., Santos F.R., Schneider M.P.C., Senanez H.N.,
RA Silva A.M.R., da Silva A.L.C., Silva D.W., Silva R., Simoes I.C.,
RA Simon D., Soares C.M.A., Soares R.B.A., Souza E.M., Souza K.R.L.,
RA Souza R.C., Steffens M.B.R., Steindel M., Teixeira S.R., Urmenyi T.,
RA Vettore A., Wassén R., Zana A., Simpson A.J.G.;
RT "The complete genome sequence of Chromobacterium violaceum reveals
RT remarkable and exploitable bacterial adaptability".
RL Proc. Natl. Acad. Sci. U.S.A. 100:11660-11665(2003).
DR EMBL; AE016920; AA060719.1; -.
DR InterPro; IPR01577; CytB561_bact.

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DR Pfam: PF01292; N1_hydr_CYTB; 1.
 KW Complete proteome
 SQ SEQUENCE 181 AA; 19650 MW; 94FBFB853494F08 CRC64;

Query Match
 Best Local Similarity 43.8%; Score 7; DB 2; Length 181;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 PLRSLG 14
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 DB 42 PLRSLG 48

RESULT 13

Q66F29 PRELIMINARY; PRT; 239 AA.
 AC Q66F29;
 DT 25-OCT-2004 (TREMBLrel. 28, Created)
 DT 25-OCT-2004 (TREMBLrel. 28, Last sequence update)
 DT 25-OCT-2004 (TREMBLrel. 28, Last annotation update)
 DE Putative ABC phosphonate transporter, ATP binding protein, also
 DE putative C-P lyase component.
 GN Name=phnL; ORFNames=YPT80511;
 OS Yersinia pseudotuberculosis IP 32953.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
 OC Enterobacteriaceae; Yersinia.
 OK NCBI_TaxID=273123;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=IP 32953;
 RX Chain F.S.G., Carmel E., Larimer F.W., Lamerdin J., Stouland P.O.,
 RA Regla W.M., Georgescu A.M., Verge L.M., Land M.L., Motin L.V.,
 RA Brubaker R.R., Fowler J., Hinnelbusch B.J., Marceau M., Medigue C.,
 RA Simonet M., Chena-Francois V., Souza B., Dacheux D., Elliott J.M.,
 RA Derise A., Hauser L.J., Garcia E.;
 RT "Insights into the genome evolution of Yersinia pestis through whole
 RT genome comparison with Yersinia pseudotuberculosis";
 RL Proc. Natl. Acad. Sci. U.S.A. 101:13826-13831 (2004).
 DR EMBL; BX936398; CAH19751.1; -;
 DR GO; GO:0016829; P:lyase activity; IEA.
 DR InterPro: IPR003593; AAA ATPase.
 DR InterPro: IPR003439; ABC_transporter.
 DR Pfam; PF00005; ABC_tran; 1.
 DR ProDom; PD000006; ABC_transporter; 1.
 DR SMART; SM00382; AAA; 1.
 DR PROSITE; PS00211; ABC_TRANSPORTER_1; 1.
 DR PROSITE; PS00893; ABC_TRANSPORTER_2; 1.
 KW ATP-binding; Lyase.
 SQ SEQUENCE 239 AA; 26695 MW; 4096220E6760B9FB CRC64;

Query Match
 Best Local Similarity 43.8%; Score 7; DB 2; Length 239;
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QY 2 QHGIRLP 8
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 DB 20 QHGIRLP 26

RESULT 14

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 AC Q82BF4; Q7CKG6;
 DT 01-MAR-2002 (TREMBLrel. 20, Created)
 DT 01-MAR-2002 (TREMBLrel. 20, Last sequence update)
 DT 25-OCT-2004 (TREMBLrel. 28, Last annotation update)
 DE Phosphonate transporter ATP-binding protein (ATP-binding component of
 DE phosphonate ABC transporter).
 GN Name=phnL; OrderedLocNames=YPO3462, Y0723;
 OS Yersinia pestis.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
 OC Enterobacteriaceae; Yersinia.

OK NCBI_TaxID=632;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CO-92 / Biovar Orientalis;
 RX MEDLINE=21470413; PubMed=11586360; DOI=10.1038/35097083;
 RA Parhill J., Wren B.W., Thomson N.R., Tibball R.W., Holden M.T.G.,
 RA Baker S., Basham D., Bentley S.D., Brooks K., Churcher C.M., Mungall K.L.,
 RA Chillingworth T., Cronin A., Davies R.M., Davis P., Dougan G.,
 RA Felwell T., Hamlin N., Holroyd S., Jagsis K., Karlyshev A.V.,
 RA Leach S., Moule S., Oyston P.C.F., Quail M.A., Rutherford K.M.,
 RA Simmonds M., Skellon J., Stevens K., Whitehead S., Barrett B.G.;
 RT "Genome sequence of Yersinia pestis, the causative agent of plague";
 RL Nature 413:523-527 (2001).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=KIMS / Biovar Mediaevalis;
 RX MEDLINE=22137863; PubMed=12142430;
 RX DOI=10.1128/JB.184.16.4601-4611.2002;
 RA Deng W., Burland V., Plunkett G. III, Boutin A., Mayhew G.F., Liss P.,
 RA Perna N.T., Rose D.J., Mau B., Zhou S., Schwartz D.C.,
 RA Fetherston J.D., Lindler L.E., Brubaker R.R., Plano G.V.,
 RA Straley S.C., McDonough K.A., Niles M.L., Mateon J.S., Blattner F.R.,
 RA Perry R.D.;
 RT "Genome sequence of Yersinia pestis KIM";
 RL J. Bacteriol. 184:4601-4611 (2002).
 CC -1- SIMILARITY: Belongs to the ABC transporter family.
 DR EMBL; AJ414157; CAC92691.1; -;
 DR EMBL; AE013674; NAM84311.1; -;
 DR PIR; AG0420; AG0420.
 DR GO; GO:0016020; C:membrane; IEA.
 DR GO; GO:0005524; F:ATP binding; IEA.
 DR GO; GO:0042626; F:ATPase activity; IEA.
 DR GO; GO:0000166; F:nucleotide binding; IEA.
 DR GO; GO:0006810; P:transport; IEA.
 DR Pfam; PF00005; ABC_tran; 1.
 DR ProDom; PD000006; ABC_transporter; 1.
 DR SMART; SM00382; AAA; 1.
 DR PROSITE; PS00211; ABC_TRANSPORTER_1; 1.
 DR PROSITE; PS00893; ABC_TRANSPORTER_2; 1.
 KW ATP-binding; Complete proteome.
 SQ SEQUENCE 239 AA; 26667 MW; 59AB94CC3760B9FB CRC64;

Query Match
 Best Local Similarity 43.8%; Score 7; DB 2; Length 239;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 QHGIRLP 8
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 DB 20 QHGIRLP 26

RESULT 15

Q9RXU2 PRELIMINARY; PRT; 259 AA.
 ID Q9RXU2;
 AC Q9RXU2;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
 DE Hypothetical protein DR0214.
 DE OrderedLocNames=DR0214;
 GN Deinococcus radiodurans.
 OS Deinococcus radiodurans.
 OC Bacteria; Deinococcus-Thermus; Deinococci; Deinococcales;
 OC Deinococcaceae; Deinococcus.
 OK NCBI_TaxID=1299;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=R1 / ATCC 13939 / DSM 20539 / NCIB 9279;
 RX MEDLINE=20036896; PubMed=10567266; DOI=10.1126/science.286.5444.1571;
 RA White O., Eisen J.A., Heidelberg J.F., Hickey E.K., Peterson J.D.,
 RA Dodson R.J., Haft D.H., Gwinn M.L., Nelson W.C., Richardson D.L.,
 RA Moffat K.S., Qin H., Jiang L., Pamphile W., Crosby M., Shen M.,
 RA Vamathavan J.J., Lam P., McDonald L.A., Uterback T.R., Zalewski C.,

RA Makarova K.S., Aravind L., Daly M.J., Minton K.W., Fleischmann R.D.,
RA Ketchum K.A., Nelson K.E., Salzberg S.L., Smith H.O., Venter J.C.,
RA Fraser C.M.;
RT "Genome sequence of the radioresistant bacterium Deinococcus
RT radiodurans R1." Science 286:1571-1577(1999).
RL EMBL; AE001883; AAF09805.1; -.
DR PIR; A75547; A75547.
DR TIGR; DR0214; -.
KW Complete proteome; Hypothetical protein.
SQ SEQUENCE 259 AA; 29103 MW; 49522C0ADD9327CF CRC64;

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Best Local Similarity 100.0%; Pred. No. 35;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 HGIRLPL 9
Db 190 HGIRLPL 196

Search completed: July 26, 2005, 16:37:18
Job time : 175 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 18:18:13 ; Search time 1937 Seconds
(without alignments)
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Title: US-10-726-967A-3

Perfect score: 16
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Post-processing: Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	517	9	AB089958 Homo sapi
2	16	100.0	1287	6	AR224122 Sequence
3	16	100.0	1287	6	AR269253 Sequence
4	16	100.0	1287	6	AR478808 Sequence

5	16	100.0	1287	6	AR487374	AR487374 Sequence
6	16	100.0	1287	6	AR532014	AR532014 Sequence
7	16	100.0	1287	6	AR540915	AR540915 Sequence
8	16	100.0	1287	6	AR560125	AR560125 Sequence
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12	16	100.0	1302	6	AR224103	AR224103 Sequence
13	16	100.0	1302	6	AR269234	AR269234 Sequence
14	16	100.0	1302	6	AR478789	AR478789 Sequence
15	16	100.0	1302	6	AR487355	AR487355 Sequence
16	16	100.0	1302	6	AR531995	AR531995 Sequence
17	16	100.0	1302	6	AR540896	AR540896 Sequence
18	16	100.0	1302	6	AR560106	AR560106 Sequence
19	16	100.0	1302	6	AX105407	AX105407 Sequence
20	16	100.0	1302	6	AX573845	AX573845 Sequence
21	16	100.0	1302	6	AX700454	AX700454 Sequence
22	16	100.0	1305	6	AR224123	AR224123 Sequence
23	16	100.0	1305	6	AR269254	AR269254 Sequence
24	16	100.0	1305	6	AR478809	AR478809 Sequence
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26	16	100.0	1305	6	AR532015	AR532015 Sequence
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ALIGNMENTS

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LOCUS	AB089958				
DEFINITION	Homo sapiens BACE mRNA for beta-site APP cleaving enzyme isoform				
VERSION	AB089958.1	GI:34014375			
KEYWORDS					
SOURCE					
ORGANISM	Homo sapiens (human)				
REFERENCE					
AUTHORS	Tanahashi, H.				
TITLE	A novel alternatively spliced isoform of BACE, I-127 induced by				
JOURNAL	cytoheximide treatment				
REFERENCE	2 (bases 1 to 517)				
AUTHORS	Tanahashi, H.				
TITLE	Direct Substitution				
JOURNAL	Submitted (17-AUG-2002) Hiroshi Tanahashi, National Institute of				
REFERENCE	Neuroscience, Division of Demyelinating Disease and Aging; 4-1-1				
AUTHORS	Ogawabashi, Kodaira, Tokyo 187-8502, Japan				
TITLE	(E-mail: tanahashicnp.go.jp, Tel: 81-042-341-2711 (ex. 5163),				
JOURNAL	Fax: 81-042-346-1747)				
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AUTHORS	1. . 517				
TITLE	/organism="Homo sapiens"				

us-10-726-967a-3.01p2n.rge

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donor in exon 3."
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Best Local Similarity: 16.00
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Gaps: 0
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Mismatch: 0
Length: 517

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ACCESSION AR224122
VERSION AR224122.1
KEYWORDS GI:2332782
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6440698-A 50 27-AUG-2002;
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Length: 1287

RESULT 3
LOCUS AR269253
DEFINITION Sequence 50 from patent US 6500667. DNA
ACCESSION AR269253
VERSION AR269253.1
KEYWORDS GI:29700221
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6500667-A 50 31-DEC-2002;
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ACCESSION AR269253
VERSION AR269253.1
KEYWORDS GI:29700221
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6500667-A 50 31-DEC-2002;
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Gaps: 0
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Conservative: 16
Mismatch: 0
Length: 1287

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DEFINITION Sequence 50 from patent US 669671. DNA
ACCESSION AR478808
VERSION AR478808.1
KEYWORDS GI:47237528
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 669671-A 50 02-MAR-2004;
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DB: 100.00%
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Gaps: 0
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Conservative: 16
Mismatch: 0
Length: 1287

RESULT 5
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DEFINITION Sequence 50 from patent US 6706485. DNA
ACCESSION AR487374
VERSION AR487374.1
KEYWORDS GI:14444444
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6706485-A 50 01-MAY-2004;
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ACCESSION AR487374
VERSION AR487374.1 GI:47252472
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Method of identifying agents that inhibit APP processing activity
JOURNAL Patent: US 6706485-A 50 16-MAR-2004;
FEATURES
source
1..1287
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3.27e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR487374 (1-1287)

QY
1 Thrglnhlglylleargleuproleuargserglyleuglygla 16
|||||
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

Db
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

RESULT 6
AR532014
LOCUS AR532014 1287 bp DNA linear PAT 08-OCT-2004
DEFINITION Sequence 50 from patent US 6727074.
ACCESSION AR532014
VERSION AR532014.1 GI:53920548
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6727074-A 50 27-APR-2004;
FEATURES
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/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3.27e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR532014 (1-1287)

QY
1 Thrglnhlglylleargleuproleuargserglyleuglygla 16
|||||
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

Db
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

RESULT 7
AR540915
LOCUS AR540915 1287 bp DNA linear PAT 08-OCT-2004
DEFINITION Sequence 50 from patent US 6737510.
ACCESSION AR540915
VERSION AR540915.1 GI:53932428

KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6737510-A 50 18-MAY-2004;
FEATURES
source
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/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3.27e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR540915 (1-1287)

QY
1 Thrglnhlglylleargleuproleuargserglyleuglygla 16
|||||
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

Db
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

RESULT 8
AR560125
LOCUS AR560125 1287 bp DNA linear PAT 08-OCT-2004
DEFINITION Sequence 50 from patent US 6753163.
ACCESSION AR560125
VERSION AR560125.1 GI:53970492
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1287)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses
JOURNAL Patent: US 6753163-A 50 22-JUN-2004;
FEATURES
source
1..1287
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3.27e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR560125 (1-1287)

QY
1 Thrglnhlglylleargleuproleuargserglyleuglygla 16
|||||
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

Db
64 ACCGACGACGGCATCCGGCTGCCCTGCCGACGCGGCTGGGGGCGCC 111

RESULT 9
AX105432
LOCUS AX105432 1287 bp DNA linear PAT 30-APR-2001
DEFINITION Sequence 50 from Patent WO0123533.
ACCESSION AX105432
VERSION AX105432.1 GI:13921541
KEYWORDS

SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE other sequences; artificial sequences.
AUTHORS 1 Gurney, M. and Bienkowski, M.J.
TITLE Alzheimer's disease secretase, app substrates therefor, and uses therefor
JOURNAL Patent: WO 0123533-A 50 05-APR-2001;
Pharmacia & Upjohn Company (US)
FEATURES
source
1..1287
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Hu-Ap2(b) delta TM"

ORIGIN
Alignment Scores:
Pred. No.: 3.27e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967a-3 (1-16) x AX573870 (1-1287)

QY 1 ThGlnHsglyIleArgLeuProLeuArgSerGlyLeuGlyGlyAla 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTCGCGCAGCGCCTGGGGGCGCC 111

RESULT 11
BD235897 1302 bp DNA linear PAT 17-JUL-2003
LOCUS Alzheimer's disease secretase.
DEFINITION

ACCESSION BD235897
VERSION BD235897.1 GI:33045667
KEYWORDS JP 2002526081-A/13.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS 1 (bases 1 to 1302)
Gurney, M.E., Bienkowski, M.J., Heinrichson, R.L., Parodi, L.A. and Yan, R.
TITLE Alzheimer's disease secretase
JOURNAL Patent: JP 2002526081-A 13 20-AUG-2002;
PHARMACIA AND UPJOHN CO
COMMENT OS Homo sapiens (human)
PN JP 2002526081-A/13
PD 20-AUG-2002
PF 24-SEP-1999 JP 2000574268
PR 24-SEP-1998 US 60/101594
PI MARK E GURNEY, MICHAEL JEROME BIENKOWSKI, ROBERT LEROY PI
HEINRICHSON, PI
LUI S A PARODI, RIOIANG YAN
PC C12N15/09, A61K45/00, A61P25/28, C07K14/47, C07K16/18, C12N1/15, PC
C12N1/19, PC
C12N1/21, C12N5/10, C12N9/64, C12P21/02, C12P21/08, C12Q1/37, G01N33/15,
PC G01N33/50// (C12N1/21, C12R1:19), C12N15/00, C12N5/00 CC
Alzheimer's disease secretase
FH Key location/Qualifiers
FT source 1..1302
/organism="Homo sapiens (human)".
location/Qualifiers
1..1302
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"

ORIGIN
Alignment Scores:
Pred. No.: 3.3e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967a-3 (1-16) x BD235897 (1-1302)

QY 1 ThGlnHsglyIleArgLeuProLeuArgSerGlyLeuGlyGlyAla 16
Db 4 ACTCAGCATGGATTCGTCCTCCACCTGCGGTGCGGTGGGTGCT 51

RESULT 12
AR224103 1302 bp DNA linear PAT 26-SEP-2002
LOCUS Sequence 25 from patent US 6440698.
DEFINITION AR224103
ACCESSION AR224103
VERSION AR224103.1 GI:23332763
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1302)
Gurney, M.E., Bienkowski, M.J., Heinrichson, R.L., Parodi, L.A. and Yan, R.
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses therefor
JOURNAL Patent: US 6440698-A 25 27-AUG-2002;
FEATURES
source
1..1302
/organism="unknown"
/mol_type="genomic DNA"

Alignment Scores:

Pred. No.: 3.3e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR224103 (1-1302)

Qy 1 ThrGlnHISGlyIleArgLeuProLeuArgSerGlyLeuGlyAla 16
|||||
Db 4 ACTCAGCATGTATTGCTCTGCCACTGCGGTGCTGGTGGTCT 51

RESULT 13 AR269234 1302 bp DNA linear PAT 10-APR-2003
LOCUS AR269234
DEFINITION Sequence 25 from patent US 6500667.
ACCESSION AR269234
VERSION AR269234.1 GI:29700202
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 1302)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and Yan,R.
TITLE Aspartyl protease 2 (Asp2) antisense oligonucleotides
JOURNAL Patent: US 6500667-A 25 31-DEC-2002;
FEATURES Location/Qualifiers
source 1..1302
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Alignment Scores:
Pred. No.: 3.3e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR269234 (1-1302)

Qy 1 ThrGlnHISGlyIleArgLeuProLeuArgSerGlyLeuGlyAla 16
|||||
Db 4 ACTCAGCATGTATTGCTCTGCCACTGCGGTGCTGGTGGTCT 51

RESULT 14 AR478789 1302 bp DNA linear PAT 14-MAY-2004
LOCUS AR478789
DEFINITION Sequence 25 from patent US 6699671.
ACCESSION AR478789
VERSION AR478789.1 GI:47237509
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 1302)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and Yan,R.

TITLE Alzheimer's disease secretase, APP substrates therefor, and uses therefor

JOURNAL Patent: US 6699671-A 25 02-MAR-2004;
FEATURES Location/Qualifiers
source 1..1302
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Alignment Scores:

Pred. No.: 3.3e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR478789 (1-1302)

Qy 1 ThrGlnHISGlyIleArgLeuProLeuArgSerGlyLeuGlyAla 16
|||||
Db 4 ACTCAGCATGTATTGCTCTGCCACTGCGGTGCTGGTGGTCT 51

RESULT 15 AR487355 1302 bp DNA linear PAT 14-MAY-2004
LOCUS AR487355
DEFINITION Sequence 25 from patent US 6706485.
ACCESSION AR487355
VERSION AR487355.1 GI:47252453
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 1302)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and Yan,R.
TITLE Method of identifying agents that inhibit APP processing activity
JOURNAL Patent: US 6706485-A 25 16-MAR-2004;
FEATURES Location/Qualifiers
source 1..1302
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Alignment Scores:
Pred. No.: 3.3e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-3 (1-16) x AR487355 (1-1302)

Qy 1 ThrGlnHISGlyIleArgLeuProLeuArgSerGlyLeuGlyAla 16
|||||
Db 4 ACTCAGCATGTATTGCTCTGCCACTGCGGTGCTGGTGGTCT 51

Search completed: July 27, 2005, 19:43:12
Job time : 1938 secs

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OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 17:25:16 ; Search time 432 Seconds
(without alignments)
219.250 Million cell updates/sec

Title: US-10-726-967A-3
Perfect score: 16
Sequence: 1 TQHGRLRLRSLGGA 16

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Ygapop 60.0 , Ygapext 60.0
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4390206 seqs, 2959870667 residues

Word size: 1

Total number of hits satisfying chosen parameters: 8770599

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

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-IOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi
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-OUTFMT=plco -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
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-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

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3: geneseqn20008:*
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12: geneseqn20048:*
13: geneseqn20048:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	1287	4	AAD17895 Human-Asp
2	16	100.0	1287	4	AAD13276 Human-Asp
3	16	100.0	1287	4	AAD06768 Human-Asp
4	16	100.0	1287	4	AAS11547 Human-CDN
5	16	100.0	1287	6	ABL52487 Human-Asp

6	16	100.0	1287	12	ADJ94362 Human-pro
7	16	100.0	1287	12	AD050458 Human-Asp
8	16	100.0	1287	13	ADR75371 Human-Asp
9	16	100.0	1302	3	AA15670 Human-pro
10	16	100.0	1302	4	AAS11713 DNA encod
11	16	100.0	1302	4	AAD17876 Human-pro
12	16	100.0	1302	4	AAD13032 Human-pro
13	16	100.0	1302	4	AAD06750 Human-pro
14	16	100.0	1302	4	AAS11528 Human-CDN
15	16	100.0	1302	6	ABL52468 Human-pro
16	16	100.0	1302	12	ADJ94337 Human-pro
17	16	100.0	1302	12	AD050433 Human-pro
18	16	100.0	1302	13	ADR75346 Human-pro
19	16	100.0	1305	4	AAS11733 DNA encod
20	16	100.0	1305	4	AAD17896 Human-Asp
21	16	100.0	1305	4	AAD13277 Human-Asp
22	16	100.0	1305	4	AAD06769 Human-Asp
23	16	100.0	1305	4	AAS11548 Human-CDN
24	16	100.0	1305	6	ABL52488 Human-Asp
25	16	100.0	1305	12	ADJ94364 Human-pro
26	16	100.0	1305	12	AD050460 Human-Asp
27	16	100.0	1305	13	ADR75373 Human-Asp
28	16	100.0	1341	3	AA15668 T7-caspase
29	16	100.0	1341	4	AAS11711 DNA encod
30	16	100.0	1341	4	AAD17874 T7-Human-
31	16	100.0	1341	4	AAD13030 T7-Human-
32	16	100.0	1341	4	AAD06748 T7-Human-
33	16	100.0	1341	4	AAS11526 Human-CDN
34	16	100.0	1341	6	ABL52466 Human-CDN
35	16	100.0	1341	12	ADJ94333 Human-CDN
36	16	100.0	1341	12	AD050429 T7-Human-
37	16	100.0	1341	13	ADR75342 T7-Human-
38	16	100.0	1362	3	AA15688 Modified
39	16	100.0	1362	3	AAS11715 DNA encod
40	16	100.0	1362	4	AAD17878 Human-Asp
41	16	100.0	1362	4	AAD13034 Human-Asp
42	16	100.0	1362	4	AAD06752 Human-Asp
43	16	100.0	1362	4	AAS11530 Human-CDN
44	16	100.0	1362	6	ABL52470 Human-Asp
45	16	100.0	1362	12	ADJ94341 Human-pro

ALIGNMENTS

RESULT 1	
AD17895	
ID	AAD17895 standard; cDNA; 1287 BP.
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AC	AAD17895;
XX	
DT	10-DEC-2001 (first entry)
XX	
DE	Human-Asp 2(b) protein lacking transmembrane domain encoding cDNA.
XX	
KW	Human; aspartyl protease 2b; Asp2b; amyloid precursor protein; APP;
KW	Alzheimer's disease; AD; dementia; neurofibrillary tangle; gliosis;
KW	amyloid plaque; neuronal loss; proteolytic; neuroprotective;
KW	ss.
XX	
OS	Homo sapiens.
OS	Synthetic.
XX	
FT	Key
FT	Location/Qualifiers
FT	1..1287
FT	/tag= a
FT	/product= "Human-Asp 2(b) protein lacking transmembrane
FT	domain"
XX	
PN	GB2357767-A.
XX	
PD	04-JUL-2001.
XX	
PF	22-SEP-2000; 2000GB-00023315.

XX 23-SEP-1999; 99US-00404133.
PR 23-SEP-1999; 99US-0155483P.
PR 23-SEP-1999; 99MO-US020081.
PR 13-OCT-1999; 99US-00416901.
PR 06-DEC-1999; 99US-0169232P.
XX
PA (PHAA) PHARMACIA & UPJOHN CO.
XX
PI Bienkowiecki MJ, Gurney M;
XX
DR WPI; 2001-444208/48.
DR P-PSDB; AAE10646.
XX
PT Polypeptide comprising fragments of human aspartyl protease with amyloid
PT precursor protein processing activity and alpha-secretase activity, for
XX identifying modulators useful in treating Alzheimer's disease.
XX
PS Example 10; Page 137, 187pp; English.
XX
XX The patent discloses human aspartyl protease 1 (hu-Asp1) or modified Asp1
CC proteins which lack transmembrane domain or amino terminal domain or
CC cytoplasmic domain and retains alpha-secretase activity and amyloid
CC protein precursor (APP) processing activity. The proteins of the
CC invention are useful for assaying hu-Asp1 alpha-secretase activity, which
CC in turn is useful for identifying modulators of hu-Asp1 alpha-secretase
CC activity, where modulators that increase hu-Asp1 alpha-secretase activity
CC are useful for treating Alzheimer's disease (AD) which causes progressive
CC dementia with consequent formation of amyloid plaques, neurofibrillary
CC tangles, gliosis and neuronal loss. Hu-Asp1 protease substrate is useful
CC for assaying hu-Asp1 proteolytic activity, by contacting hu-Asp1 protein
CC with the substrate under acidic conditions and determining the level of
CC hu-Asp1 proteolytic activity. The present sequence is a cDNA encoding
CC human Asp 2(b) protein lacking a transmembrane (TM) domain which is
CC generated by the deletion of the C-terminal TM domain and intracellular
XX domains of human Asp 2(b) protein
XX
SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;
XX
Alignment Scores:
Pred. No.: 2,64e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0
DB: 4
US-10-726-967A-3 (1-16) x AAD17895 (1-1287)
OY 1 Thrglnhsglylyleargyleuproleuhrgserglyleuglyyala 16
Db 64 ACCGAGCAGCGGCAATCCGGCTCCCTGCGCAGCGGCGGCGGCGCC 111
RESULT 2
ID AAD13276 standard; cDNA; 1287 BP.
XX
AC AAD13276;
XX
DT 23-OCT-2001 (first entry)
XX
DE Human-Asp2(b) deltaTM protein cDNA.
XX
XX Human; aspartyl protease 2b; Asp 2b; beta-amyloid precursor protein; APP;
KM beta-secretase; Alzheimer's disease; dementia; amyloid plaque; gliosis;
KM neurofibrillary tangle; neuronal loss; amyloid-beta peptide; nootropic;
KM neuroprotective; antisense therapy; Asp2(b) deltaTM protein;
KM gene therapy; ss.
XX
XX Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers

FT CDS 1..1287
FT /*tag= a
FT /product= "Human Asp2 (b) deltaTM protein"
XX
XX MO200150829-A2.
XX
XX 19-JUL-2001.
XX
XX 09-MAY-2001; 2001WO-1B000799.
XX
XX 09-MAY-2001; 2001WO-1B000799.
XX
XX 09-MAY-2001; 2001WO-1B000799.
XX
XX (BIEN/) BIENKOWSKI M J.
PA (GURN/) GURNEY M E.
PA (HEIN/) HEINRIKSON R L.
PA (PARO/) PARODI L A.
XX (YANR/) YAN R.
XX
XX Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
PI WPI; 2001-483072/52.
XX
XX P-PSDB; AAE06891.
XX
XX Novel purified polypeptide comprising fragment of mammalian aspartyl
PT protease 2, lacking Asp2 transmembrane domain and retaining beta
PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
XX activity.
XX
XX Example 10; Page 166-167; 185pp; English.
XX
XX The invention relates to human aspartyl proteases (Hu-Asp), beta-amyloid
XX precursor protein (APP) isoforms and their corresponding DNA molecules.
XX Human aspartyl proteases can act as beta-secretase proteases useful for
XX treating Alzheimer's disease. APP isoforms are useful for identifying
XX modulators of amyloid-beta peptide production, for use in designing
XX therapeutics for the treatment and prevention of Alzheimer's disease,
XX dementia, formation of amyloid plaques, neurofibrillary tangles, gliosis
XX and neuronal loss. APP isoforms are also used in methods for identifying
XX inhibitors and modulators of human Asp2 activity. The invention relates
XX to a method for identifying agents that modulate the activity of human
XX aspartyl protease Asp2. Amyloid-beta peptides obtained from APP are used
XX as a means to screen in cellular assays for the inhibitors of beta- and
XX gamma- secretase. Hu-Asp DNA fragments are useful as probes or primers in
XX polymerase chain reactions (PCR). The probes are useful for detecting Hu-
XX Asp nucleic acids in in vitro assays and in Northern and Southern blots.
XX The present cDNA sequence encodes Human aspartyl protease 2b (Hu-Asp2b)
XX deltaTM protein which is obtained by the deletion of C-terminal
XX transmembrane and intracellular domains of Hu-Asp2b. Human Asp2b has beta
XX -secretase activity
XX
SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;
XX
XX
Alignment Scores:
Pred. No.: 2,64e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0
DB: 4
US-10-726-967A-3 (1-16) x AAD13276 (1-1287)
OY 1 Thrglnhsglylyleargyleuproleuhrgserglyleuglyyala 16
Db 64 ACCGAGCAGCGGCAATCCGGCTCCCTGCGCAGCGGCGGCGGCGCC 111
RESULT 3
ID AAD06768 standard; cDNA; 1287 BP.
XX
AC AAD06768;
XX
DT 10-AUG-2001 (first entry)

```
XX Human aspartyl protease 2 (b) delta TM cDNA.
DE
XX
XX Human; alpha-secretase; amyloid precursor protein; APP; therapy;
KW Alzheimer's disease; antialzheimer's; aspartyl protease 2; Asp 2;
XX beta-secretase; chromosome 11q23.3-24.1; mutant; ss.
XX
OS Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
XX CDS 1..1287
XX /tag= a
XX /product= "Human aspartyl protease 2 (b) delta TM"
XX
XX WO200123533-A2.
XX
XX 05-APR-2001.
XX
XX 22-SEP-2000; 2000WO-US026080.
XX
XX 23-SEP-1999; 99US-0155493P.
XX 23-SEP-1999; 99WO-US020881.
XX 13-OCT-1999; 99US-00416901.
XX 06-DEC-1999; 99US-0169232P.
XX
XX (PHAA ) PHARMACIA & UPJOHN CO.
XX
XX Gurney M, Bienkowski MJ;
XX
XX WPI: 2001-290516/30.
XX P-PSDB; AAE02598.
XX
XX Enzymes that cleave the alpha-secretase site of the amyloid precursor
XX protein, useful for the treatment of Alzheimer's disease.
XX
XX Example 10; Page 165-166; 189pp; English.
XX
XX The present invention relates to enzymes for cleaving the alpha-
XX secretase site of the amyloid precursor protein (APP) and methods of
XX identifying those enzymes. The methods may be used to identify enzymes
XX that may be used to cleave the alpha-secretase cleavage site of the APP
XX protein. The enzymes may be used to treat or modulate the progress of
XX Alzheimer's disease. The present sequence is human aspartyl protease 2
XX (Asp 2) (b) delta TM cDNA. The Asp 2 gene from which it is derived is
XX located on chromosome 11q23.3-24.1. The Asp 2 has beta-secretase protease
XX activity
XX
XX Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;
XX
XX Alignment Scores:
XX Pred. No.: 2.64e-06 Length: 1287
XX Score: 16.00 Matches: 16
XX Percent Similarity: 100.00% Conservative: 0
XX Best Local Similarity: 100.00% Mismatches: 0
XX Query Match: 100.00% Indels: 0
XX DB: 4 Gaps: 0
XX
XX US-10-726-967A-3 (1-16) x AAD06768 (1-1287)
XX
XX 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyIleGlyAla 16
XX |||||
XX 64 ACCGACGACGGCATCCGGCTGCCCTGCGACGGCGCTGGGAGCGCC 111
XX
XX RESULT 4
XX AAS11547
XX ID AAS11547 standard; cDNA; 1287 BP.
XX
XX AAS11547;
XX
XX 24-OCT-2001 (first entry)
XX
XX Human cDNA encoding Human-pro-Asp 2 (b) delta TM.
```

```
XX Human; Aspartyl protease; beta-secretase; nontropic; ASP2;
KW neuroprotective; amyloid protein precursor; APP; Alzheimer's disease;
XX amyloid-beta; Abeta; Human-pro-Asp 2(b) delta TM; ss; mutant.
XX
OS Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
XX CDS 1..1287
XX /tag= a
XX /product= "Human-Pro-Asp 2(b) delta TM"
XX
XX WO200149098-A2.
XX
XX 12-JUL-2001.
XX
XX 09-MAY-2001; 2001WO-IB000798.
XX
XX 09-MAY-2001; 2001WO-IB000798.
XX
XX (BIEN/) BIENKOWSKI M J.
XX (GURN/) GURNEY M E.
XX (HEIN/) HEINRIKSON R L.
XX (PARO/) PARODI L A.
XX (YANR/) YAN R.
XX
XX Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
XX
XX WPI: 2001-502549/55.
XX
XX Novel purified polypeptide comprising fragment of mammalian aspartyl
XX protease 2, lacking Asp2 transmembrane domain and retaining beta
XX secretase activity of Asp2 useful for identifying inhibitors of Asp2
XX activity.
XX
XX Disclosure; Page 166-167; 185pp; English.
XX
XX The invention relates to a purified polypeptide comprising a fragment of
XX mammalian aspartyl protease (Asp) 2 protein which lacks the Asp2
XX transmembrane domain and the Asp2 protein, and where the polypeptide and
XX the fragment retain the beta-secretase activity of the mammalian Asp2
XX protein. The invention also details polynucleotides for the Asp proteins
XX and vectors expressing them, and a polypeptide (isoform of amyloid
XX protein precursor (APP) comprising the amino acid sequence of an APP or
XX its fragment containing an APP cleavage site recognizable by a mammalian
XX beta-secretase, and further comprising two lysine residues at the
XX carboxyl terminus of the amino acid sequence of the mammalian APP or APP
XX fragment. Also included in the invention are methods of identifying
XX modulators or inhibitors of Asp2. Modulators and inhibitors of Asp2 are
XX useful for treating Alzheimer's disease. APP is useful in methods for
XX identifying inhibitors or modulators of human Asp2 activity and amyloid-
XX beta (Abeta) peptide production. APP is also useful in designing
XX therapeutics for the treatment or prevention of Alzheimer's disease. APP
XX comprising the APP-Sw-beta-secretase peptide sequence (NLDA), which is
XX associated with increased levels of Abeta processing is useful in assays
XX relating the Alzheimer's research. The expression vector is useful for
XX recombinantly expressing APP. Nucleic acids that hybridize to Asp
XX oligonucleotides are useful as probes or primers. The probes are useful
XX for detecting Hu-Asp nucleic acids in in vitro assays and in Northern and
XX Southern blots. The present sequence encodes Human-pro-Asp 2(b) delta TM
XX protein, which lacks the C-terminal transmembrane domain
XX
XX Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;
XX
XX Alignment Scores:
XX Pred. No.: 2.64e-06 Length: 1287
XX Score: 16.00 Matches: 16
XX Percent Similarity: 100.00% Conservative: 0
XX Best Local Similarity: 100.00% Mismatches: 0
XX Query Match: 100.00% Indels: 0
XX DB: 4 Gaps: 0
```


CC antibody that specifically binds to Hu-Asp polypeptides, identifying a
CC cell that can be used to screen for inhibitors of beta secretase
CC activity, novel isoforms of amyloid protein precursor (APP), where the
CC last 2 carboxy terminus amino acids of that isoform are both lysine
CC residues (e.g. those designated APP695-KK or carrying the Swedish
CC mutation where KM at 595-596 is mutated to NL, designated e.g. APP695-SW
CC or APP695-SW-KK, or a V to F mutation at 642, e.g. APP695-VF, all useful
CC for assaying for beta secretase activity and screening for inhibitors of
CC beta-secretase) and polynucleotides that encode the APP proteins. The
CC method is useful for identifying agents that modulate the activity
CC (amyloid precursor protein processing activity) of Asp2 aspartyl
CC protease. Preferably, the method is useful for identifying agents that
CC inhibit Asp2 aspartyl protease activity. The inhibitors of amyloid
CC precursor protein processing, are useful for treating or preventing
CC Alzheimer's disease. The present sequence encodes an aspartyl protease
CC mutant construct (e.g. lacking a transmembrane domain and/or including a
CC caspase cleavage site) used to investigate the cleavage activity of Asp2
CC proteins.

XX SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 2,64e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 12 Gaps: 0

US-10-726-967A-3 (1-16) x ADJ94362 (1-1287)

QY 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyAla 16
|||||
DB 64 ACCCAGCAGCGCATCCGCTGCCCTGCGCAGCGCTCGGGCGGCC 111

RESULT 7
ADOS0458
ID ADOS0458 standard; DNA; 1287 BP.
XX
AC ADOS0458;
XX
DT 29-JUL-2004 (first entry)
XX
DE Human Asp2(b)deltaTM mutant DNA.
XX
KW Aspartyl protease; Asp; beta secretase; amyloid precursor protein; APP;
KM Alzheimer's disease; gene therapy; human; mutant; gene; de.
XX
OS Homo sapiens.
OS Synthetic.
FH
FT Key Location/Qualifiers
FT CDS 1..1287
FT /tag= a
FT /product= "Human Asp2(b) mutant protein"

XX PN US6737510-B1.
XX PD 18-MAY-2004.
XX PF 12-APR-2000; 2000US-00548373.
XX PR 24-SEP-1998; 98US-0101594P.
XX PR 23-SEP-1999; 99US-00404133.
XX PR 23-SEP-1999; 99US-0155493P.
XX PR 23-SEP-1999; 99MO-US020881.
XX PR 13-OCT-1999; 99US-00416901.
XX
XX (PHAA) PHARMACIA & UPJOHN CO.
XX PA
XX Gurney ME, Bienkowski MJ, Heinrikson RL, Parodi LA, Yan R;
XX PI
XX WPI; 2004-387112/36.

DR P-PSDB; ADO50459.
XX
XX New Asp2 aspartyl protease protein comprising tripeptides DTG and DSG
PT involved in processing amyloid precursor protein into amyloid beta,
PT useful in preparing a composition for treating or preventing Alzheimer's
PT disease.
XX
XX Example 10; SEQ ID NO 50; 108pp; English.
XX
XX The invention relates to a method for identifying an agent that decreases
CC the protease activity of the aspartyl protease (Asp) polypeptide. It also
CC provides enzyme and enzymatic procedures for cleaving the beta secretase
CC cleavage site of the amyloid precursor protein (APP). The invention is
CC useful in preparing a composition for treating or preventing Alzheimer's
CC disease. It is also useful in gene therapy. The present sequence is human
CC Asp2(b) mutant DNA. This sequence is used to illustrate the method of the
XX invention.

XX SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 2,64e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 12 Gaps: 0

US-10-726-967A-3 (1-16) x ADO50458 (1-1287)

QY 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyAla 16
|||||
DB 64 ACCCAGCAGCGCATCCGCTGCCCTGCGCAGCGCTCGGGCGGCC 111

RESULT 8
ADR75371
ID ADR75371 standard; DNA; 1287 BP.
XX
AC ADR75371;
XX
DT 18-NOV-2004 (first entry)
XX
DE Human Asp2(b)deltaTM mutant DNA.
XX
KW Aspartyl protease; Asp; amyloid precursor protein; APP; amyloid beta;
KM chromosome identification; Alzheimer's disease; human; mutant; gene; de.
XX
OS Homo sapiens.
OS Synthetic.
FH
FT Key Location/Qualifiers
FT CDS 1..1287
FT /tag= a
FT /product= "Human Asp2(b) mutant protein"

XX PN US2004166507-A1.
XX PD 26-AUG-2004.
XX PF 29-AUG-2003; 2003US-00652045.
XX PR 24-SEP-1998; 98US-0101594P.
XX PR 23-SEP-1999; 99US-00404133.
XX PR 23-SEP-1999; 99US-0155493P.
XX PR 13-OCT-1999; 99US-00416901.
XX
XX (GURN/) GURNEY M E.
XX (BIEN/) BIENKOWSKI M J.
XX (HEIN/) HEINRIKSON R L.
XX (PARO/) PARODI L A.
XX (YANR/) YAN R.
XX
XX Gurney ME, Bienkowski MJ, Heinrikson RL, Parodi LA, Yan R;
XX PI

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XX WPI; 2004-624916/60.
DR P-PSDB; ADR75372.
XX
PT Novel purified/isolated polynucleotide encoding polypeptide having
PT aspartyl protease activity involved in processing amyloid precursor
PT protein into amyloid beta, useful in identifying agent decreasing
PT activity of aspartyl protease.
XX
XX Example 10; SEQ ID NO 50; 107pp; English.
XX
CC The invention relates to nucleic acid sequences encoding aspartyl
CC protease (Asp) polypeptides having aspartyl protease activity involved in
CC processing amyloid precursor protein (APP) into amyloid beta. The
CC invention also relates to a method for identifying an agent that
CC decreases the protease activity of the Asp. Asp DNA is useful in
CC chromosome identification as they can hybridize with a specific location
CC on a human chromosome and in identifying the relationship between genes
CC and diseases (particular gene responsible for causing diseases). It is
CC also useful for identifying candidates to modulate the progression of
CC Alzheimer's disease. Asp is useful in raising antibodies that are useful
CC in diagnostic assay for detecting Hu-Asp polypeptide expression. The
CC present sequence is the human Asp2(b) deltaTM mutant DNA. This sequence is
CC used to illustrate the method of the invention.
XX
SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;
XX
Alignment Scores:
Pred. No.: 2.64e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 13 Gaps: 0
XX
US-10-726-967A-3 (1-16) x ADR75371 (1-1287)
XX
OY 1 ThGlnHieGlyTlleArgLeuProLeuAArgSerGlyLeuGlyGlyAla 16
Db 64 ACCACACACGCGATCCGCGCTGCCCTGCGACGCGCTGGGGGCGCC 111
XX
RESULT 9
AA15670
ID AA15670 standard; DNA; 1302 BP.
XX
AC AA15670;
XX
DT 15-SEP-2003 (revised)
DT 06-AUG-2003 (revised)
DT 03-AUG-2000 (first entry)
XX
DE Human-Pro-Asp-2(a)-deltaTM nucleotide sequence.
XX
KW Aspartyl protease; aspartase; amyloid precursor protein; APP; Asp 2; ss;
KW Alzheimer's disease; beta secretase site; human-pro-Asp-2(a)-deltaTM.
XX
OS Homo sapiens.
OS Enterobacteria phage T7.
OS Chimeric.
XX
XX WO200017369-A2.
XX
XX 30-MAR-2000.
XX
XX 23-SEP-1999; 99WO-US020881.
XX
XX 24-SEP-1998; 98US-0101594P.
XX
XX (PhA) PHARMACIA & UPJOHN CO.
XX
XX Gurney ME, Bienkowski MJ, Heintikson RL, Parodi LA, Yan R;
XX WPI; 2000-303209/26.
XX

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DR P-PSDB; AAY88433.
XX
XX New enzyme designated human aspartase useful in research into Alzheimer's
PT Disease is capable of cleaving amyloid protein precursor at the beta
PT secretase site to produce amyloid beta peptide.
XX
XX Example 9; Fig 8; 183pp; English.
XX
CC This sequence represents a modified version of the human aspartase 2
CC (Asp2) nucleotide sequence. The sequence is used in the bacterial
CC expression of human Asp2L. The invention relates to a protease (e.g.,
CC Asp2) capable of cleaving the beta secretase site of amyloid precursor
CC protein (APP). The protease contains a sequence encoding the amino acid
CC sequence DNG and a sequence encoding DSG or DNG separated by 100-300
CC amino acids. When mutated the Asp gene causes an autosomal dominant form
CC of Alzheimer's disease. APP localizes to the cell surface membrane and
CC have a single C-terminal transmembrane domain. Proteolytic processing of
CC APP produces the amyloid beta protein, which is possibly very important
CC in Alzheimer's disease. The invention includes a nucleotide sequence
CC encoding the protease, a vector containing the nucleotide sequence, and a
CC cell line comprising the vector. Methods for screening for inhibitors of
CC aspartase activity are also given in the invention. The human
CC aspartase protein and nucleotide sequences and the methods for
CC identifying inhibitors of the protease, are useful in the treatment of
CC and research in to Alzheimer's disease. (Updated on 06-AUG-2003 to
CC correct OS field.) (Updated on 15-SEP-2003 to standardise OS field)
XX
SQ Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;
XX
Alignment Scores:
Pred. No.: 2.67e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0
XX
US-10-726-967A-3 (1-16) x AA15670 (1-1302)
XX
OY 1 ThGlnHieGlyTlleArgLeuProLeuAArgSerGlyLeuGlyGlyAla 16
Db 4 ACTCAGCATGCTATTCGTCGCACTGCGTAGCGGCTCGGGTGGTCT 51
XX
RESULT 10
AA11713
ID AA11713 standard; DNA; 1302 BP.
XX
AC AA11713;
XX
DT 11-SEP-2003 (revised)
DT 24-OCT-2001 (first entry)
XX
DE DNA encoding T7-human aspartyl protease 2a deltaTM (low GC).
XX
KW Human; aspartyl protease 1; Asp-1; neurotropic; neuroprotective;
KW aspartyl protease 2; Asp2; amyloid protein precursor; APP;
KW beta-secretase; Alzheimer's disease; ds.
XX
XX Homo sapiens.
XX
XX Enterobacteria phage T7.
XX
XX Key Location/Qualifiers
FT 1..1302
FT CDS /*tag= a
FT /product= "T7-Aspartyl protease-2a delta TM (low GC)"
XX
XX WO200149097-A2.
XX
XX 12-JUL-2001.
XX
XX 09-MAY-2001; 2001WO-IB000797.
XX
XX 09-MAY-2001; 2001WO-IB000797.
XX

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```

XX (BIEN/) BIENKOWSKI M J.
PA (GURNEY) GURNEY M B.
PA (HEIN/) HEINRIKSON R L.
PA (PARO/) PARODI L A.
PA (YANR/) YAN R.
PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
XX WPI; 2001-502548/55.
DR P-PSDB; AAU07113.
XX
XX Novel purified polypeptide comprising fragment of mammalian aspartyl
PT protease 2, lacking Asp2 transmembrane domain and retaining beta
PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
PT activity.
XX
XX Example 9; Fig 8; 185pp; English.
XX
XX The invention relates to a novel purified polypeptide comprising a
CC fragment of mammalian aspartyl protease 2 (Asp2) protein which lacks the
CC Asp2 transmembrane domain and the Asp2 protein, and where the polypeptide
CC and the fragment retain the beta-secretase activity of the mammalian Asp2
CC protein. Also included is an isoform of amyloid protein precursor (APP)
CC comprising the amino acid sequence of a APP or its fragment containing an
CC APP cleavage site recognisable by a mammalian beta-secretase, and further
CC comprising two lysine residues at the carboxyl terminus of the amino acid
CC sequence of the mammalian APP or APP fragment. The polypeptides are used
CC for assaying for modulators of beta-secretase activity; identifying
CC agents that inhibit the APP processing activity of human Asp2 aspartyl
CC protease (hu-Asp2); identifying agents that modulate the activity of Asp2
CC; and for reducing cellular production of amyloid beta (Abeta) from APP.
CC Agents identified by the above methods are useful for treating
CC Alzheimer's disease; for identifying modulators of amyloid-beta (Abeta)
CC peptide production; and for use in designing therapeutics for the
CC treatment or prevention of Alzheimer's disease. Probes and primers
CC derived from Asp nucleic acid sequences are useful for detecting hu-Asp
CC nucleic acids in in vitro assays and in Northern and Southern blots. The
CC present sequence represents the coding sequence of T7-human Asp-2a delta
CC TM (low GC) construct which has a T7 tag, has the GC content of the 5'
CC sequence reduced by site-directed mutagenesis, and lacks the
CC transmembrane domain. This construct was used for bacterial expression
CC and purification of human Asp2a. (Updated on 11-SEP-2003 to standardise
CC OS field)
XX
XX SQ Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;
XX
XX Alignment Scores:
XX Pred. No.: 2.67e-06 Length: 1302
XX Score: 16.00 Matches: 16
XX Percent Similarity: 100.00% Conservative: 0
XX Best Local Similarity: 100.00% Mismatches: 0
XX Query Match: 100.00% Indels: 0
XX DB: 4 Gaps: 0
XX
XX US-10-726-967A-3 (1-16) x AAS11713 (1-1302)
XX
XX QY 1 ThrGlnHisGlyTlleArgLeuProLeuArgSerGlyLeuGlyVala 16
XX |||||
XX Db 4 ACTCAGCATGATGTTCTGTCGCCACTGCGGATCGGTGATGCT 51
XX
XX RESULT 11
XX AAD17876
XX AAD17876 standard; cDNA; 1302 BP.
XX
XX AC AAD17876;
XX
XX 10-DEC-2001 (first entry)
XX
XX Human-pro-Asp 2(a) protein lacking TM domain (low GC) encoding cDNA.
XX
XX Human: aspartyl protease 1; Asp1; amyloid precursor protein; APP;
XX Alzheimer's disease; AD; dementia; neurofibrillary tangle; gliosis;
XX

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```

KW amyloid plaque; neuronal loss; proteolytic; nootropic; neuroprotective;
KW Human-pro-Asp 2(a) protein; ss.
XX
XX OS Homo sapiens.
XX OS Synthetic.
XX
XX Key Location/Qualifiers
XX CDS 1..1302
XX FT /*tag= a
XX FT /product= "Human-pro-Asp 2(a) protein lacking
XX FT transmembrane domain"
XX
XX GB2357767-A.
XX
XX PN 04-JUL-2001.
XX
XX PF 22-SEP-2000; 2000GB-00023315.
XX
XX PR 23-SEP-1999; 99US-00404133.
XX PR 23-SEP-1999; 99US-0155493P.
XX PR 23-SEP-1999; 99WO-US020881.
XX PR 13-OCT-1999; 99US-00416901.
XX PR 06-DEC-1999; 99US-0169232P.
XX
XX (PHAA ) PHARMACIA & UPJOHN CO.
XX
XX PI Bienkowski MJ, Gurney M;
XX
XX WPI; 2001-444208/48.
XX
XX DR P-PSDB; AAE10640.
XX
XX PT Polypeptide comprising fragments of human aspartyl protease with amyloid
XX precursor protein processing activity and alpha-secretase activity, for
XX identifying modulators useful in treating Alzheimer's disease.
XX
XX Example 9; Fig 8; 187pp; English.
XX
XX PS The patent discloses human aspartyl protease 1 (hu-Asp1) or modified Asp1
XX proteins which lack transmembrane domain or amino terminal domain or
XX cytoplasmic domain and retains alpha-secretase activity and amyloid
XX protein precursor (APP) processing activity. The proteins of the
XX invention are useful for assaying hu-Asp1 alpha-secretase activity, which
XX in turn is useful for identifying modulators of hu-Asp1 alpha-secretase
XX activity, where modulators that increase hu-Asp1 alpha-secretase activity
XX are useful for treating Alzheimer's disease (AD) which causes progressive
XX dementia with consequent formation of amyloid plaques, neurofibrillary
XX tangles, gliosis and neuronal loss. Hu-Asp1 protease substrate is useful
XX for assaying hu-Asp1 proteolytic activity, by contacting hu-Asp1 protein
XX with the substrate under acidic conditions and determining the level of
XX hu-Asp1 proteolytic activity. The present sequence is a cDNA encoding
XX human-pro-Asp 2(a) protein lacking a transmembrane (TM) domain (low GC)
XX which is generated from human Asp 2(a) protein by the deletion of its C-
XX terminal transmembrane domain and change of degenerate codons bases in 15
XX amino acid positions from G/C to A/T to reduce the GC content
XX
XX SQ Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;
XX
XX Alignment Scores:
XX Pred. No.: 2.67e-06 Length: 1302
XX Score: 16.00 Matches: 16
XX Percent Similarity: 100.00% Conservative: 0
XX Best Local Similarity: 100.00% Mismatches: 0
XX Query Match: 100.00% Indels: 0
XX DB: 4 Gaps: 0
XX
XX US-10-726-967A-3 (1-16) x AAD17876 (1-1302)
XX
XX QY 1 ThrGlnHisGlyTlleArgLeuProLeuArgSerGlyLeuGlyVala 16
XX |||||
XX Db 4 ACTCAGCATGATGTTCTGTCGCCACTGCGGATCGGTGATGCT 51
XX
XX RESULT 12
XX AAD13032
XX

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ID AAD13032 standard; cDNA; 1302 BP.
 AC AAD13032;
 DT 23-OCT-2001 (first entry)
 XX Human-pro-Asp2(a) deltatm (low GC) protein cDNA.
 DE Human; aspartyl protease 2a; Asp 2a; beta-amyloid precursor protein; APP;
 KM beta-secretase; Alzheimer's disease; dementia; amyloid plaque; gliosis;
 KM neurofibrillary tangle; neuronal loss; amyloid-beta peptide; nootropic;
 KM neuroprotective; antisense therapy; pro-Asp2(a) deltatm protein;
 KM gene therapy; ss.
 XX Homo sapiens.
 OS Synthetic.
 XX Key Location/Qualifiers
 FH 1..1302
 FT /*tag= a
 FT /product= "Human-pro-Asp2(a) deltatm (low GC) protein"
 XX WO200150829-A2.
 XX 19-JUL-2001.
 XX 09-MAY-2001; 2001WO-IB000799.
 XX 09-MAY-2001; 2001WO-IB000799.
 XX (BIEN/) BIENKOWSKI M J.
 XX (GURN/) GURNEY M E.
 XX (HEIN/) HEINRIKSON R L.
 XX (PARO/) PARODI L A.
 XX (YANR/) YAN R.
 XX PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
 XX WPI; 2001-483072/52.
 DR P-PSDB; AAE06870.
 XX Novel purified polypeptide comprising fragment of mammalian aspartyl
 PT protease 2, lacking Asp2 transmembrane domain and retaining beta
 PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
 PT activity.
 XX Example 9; Fig 8; 185pp; English.
 XX The invention relates to human aspartyl proteases (Hu-Asp), beta-amyloid
 CC precursor protein (APP) isoforms and their corresponding DNA molecules.
 CC Human aspartyl proteases can act as beta-secretase proteases useful for
 CC treating Alzheimer's disease. APP isoforms are useful for identifying
 CC modulators of amyloid-beta peptide production, for use in designing
 CC therapeutics for the treatment and prevention of Alzheimer's disease,
 CC dementia, formation of amyloid plaques, neurofibrillary tangles, gliosis
 CC and neuronal loss. APP isoforms are also used in methods for identifying
 CC inhibitors and modulators of human Asp2 activity. The invention relates
 CC to a method for identifying agents that modulate the activity of human
 CC aspartyl protease Asp2. Amyloid-beta peptides obtained from APP are used
 CC as a means to screen in cellular assays for the inhibitors of beta- and
 CC gamma- secretase. Hu-Asp DNA fragments are useful as probes or primers in
 CC polymerase chain reactions (PCR). The probes are useful for detecting Hu-
 CC Asp nucleic acids in in vitro assays and in Northern and Southern blots.
 CC The present cDNA sequence encodes human-pro-aspartyl protease 2a (Asp2a)
 CC deltatm (low GC) protein which is obtained by the deletion of C-terminal
 CC transmembrane domain and change of degenerate codons bases in 15 amino
 CC acid positions from G/C to A/T in the Hu-Asp2a. Human Asp2a has beta-
 CC secretase activity
 XX Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;
 Alignment Scores: 2.67e-06 Length: 1302
 Pred. No.:

Score: 16.00 Matches: 16
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 4 Gaps: 0
 US-10-726-967A-3 (1-16) x AAD13032 (1-1302)
 QY 1 ThGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyGlyAla 16
 Db 4 ACTCAGCATGTATTCGTCTGCCACTGCCGTAGCGGCTCGAGTGGTCT 51
 RESULT 13
 AAD06750
 ID AAD06750 standard; cDNA; 1302 BP.
 AC AAD06750;
 XX 10-AUG-2001 (first entry)
 DT Human-pro-Asp-2(a) deltatm protein cDNA.
 DE Human; alpha-secretase; amyloid precursor protein; APP; therapy;
 KM Alzheimer's disease; antialzheimer's; aspartyl protease 2a; Asp2a;
 KM beta-secretase; Asp-2a delta TM; ss.
 XX Homo sapiens.
 OS Synthetic.
 XX Key Location/Qualifiers
 FH 1..1302
 FT /*tag= a
 FT /product= "Human-pro-Asp-2(a) delta TM protein"
 FT CDS
 XX WO200123533-A2.
 XX 05-APR-2001.
 XX 22-SEP-2000; 2000WO-US026080.
 XX 23-SEP-1999; 99US-0155493P.
 XX 23-SEP-1999; 99WO-US020881.
 XX 13-OCT-1999; 99US-00416901.
 XX 06-DEC-1999; 99US-0169232P.
 XX (PHAA) PHARMACIA & UPJOHN CO.
 XX PI Gurney M, Bienkowski MJ;
 XX WPI; 2001-290516/30.
 DR P-PSDB; AAE02592.
 XX Enzymes that cleave the alpha-secretase site of the amyloid precursor
 PT protein, useful for the treatment of Alzheimer's disease.
 XX Example 9; Page 155; 189pp; English.
 XX The present invention relates to enzymes for cleaving the alpha-
 CC secretase site of the amyloid precursor protein (APP) and methods of
 CC identifying those enzymes. The methods may be used to identify enzymes
 CC that may be used to cleave the alpha-secretase cleavage site of the APP
 CC protein. The enzymes may be used to treat or modulate the progress of
 CC Alzheimer's disease. The present sequence is a cDNA encoding human
 CC aspartyl protease 2a (Asp-2a) deltatm protein which is obtained by
 CC deleting the transmembrane domain and adding a T7 tag at the N-terminal
 CC end. This sequence has beta-secretase protease activity. Note: The
 CC present sequence is also shown in figure 8 of the specification, but
 CC lacks nucleotides at its 3' end. This sequence shown in figure 8 has a
 CC stop codon at its 3' end
 XX Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;
 Alignment Scores:
 Pred. No.:

Pred. No.: 2.67e-06 Length: 1302
 Score: 16.00 Matches: 16
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x AAD06750 (1-1302)

Qy 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyVala 16
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 4 ACTGACATGATTCGTCTGCGCACTGCGTACGCGTCTGGTGTCT 51

RESULT 14
 AAS11528 ID AAS11528 standard; cDNA; 1302 BP.
 XX
 AC AAS11528;
 XX
 DT 24-OCT-2001 (first entry)
 XX
 DE Human cDNA encoding Human-pro-Asp 2(a) delta TM (low GC).
 XX
 KW Human: Aspartyl protease; beta-secretase; neurotrophic; ASP2;
 KW neuroprotective; amyloid protein precursor; APP; Alzheimer's disease;
 KW amyloid-beta; Abeta; ss; Human-pro-Asp 2(a) delta TM (low GC).
 XX
 OS Homo sapiens.
 OS Synthetic.
 OS
 FH Key Location/Qualifiers
 FT CDS 1..1302
 FT /tag= a
 FT /product= "Human-pro-Asp 2(a) delta TM (low GC)"

MO200149098-A2.
 XX
 PD 12-JUL-2001.
 XX
 PF 09-MAY-2001; 2001WO-1B000798.
 XX
 PR 09-MAY-2001; 2001WO-1B000798.
 XX
 PA (BIEN/) BIERKOWSKI M J.
 PA (GURN/) GURNEY M E.
 PA (HEIN/) HEINRIKSON R L.
 PA (PARO/) PARODI L A.
 PA (YANR/) YAN R.
 XX
 PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
 PI
 DR MPI; 2001-502549/55.
 DR P-PSDB; AAU06614.
 XX
 PT Novel purified polypeptide comprising fragment of mammalian aspartyl
 PT protease 2, lacking Asp2 transmembrane domain and retaining beta
 PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
 PT activity.
 XX
 PS Example 9; Fig 8; 185pp; English.

The invention relates to a purified polypeptide comprising a fragment of mammalian aspartyl protease (Asp) 2 protein which lacks the Asp2 transmembrane domain and the Asp2 protein, and where the polypeptide and the fragment retain the beta-secretase activity of the mammalian Asp2 protein. The invention also details polynucleotides for the Asp proteins and vectors expressing them, and a polypeptide (isoform of amyloid protein precursor (APP)) comprising the amino acid sequence of an APP or its fragment containing an APP cleavage site recognizable by a mammalian beta-secretase, and further comprising two lysine residues at the carboxyl terminus of the amino acid sequence of the mammalian APP or APP fragment. Also included in the invention are methods of identifying modulators or inhibitors of Asp2. Modulators and inhibitors of Asp2 are

CC useful for treating Alzheimer's disease. APP is useful in methods for
 CC identifying inhibitors or modulators of human Asp2 activity and amyloid-
 CC beta (Abeta) peptide production. APP is also useful in designing
 CC therapeutics for the treatment or prevention of Alzheimer's disease. APP
 CC comprising the APP-Sw-beta-secretase peptide sequence (NDA), which is
 CC associated with increased levels of Abeta processing is useful in assays
 CC relating the Alzheimer's research. The expression vector is useful for
 CC recombinantly expressing APP. Nucleic acids that hybridize to Asp
 CC oligonucleotides are useful as probes or primers. The probes are useful
 CC for detecting Hu-Asp nucleic acids in in vitro assays and in Northern and
 CC Southern blots. The present sequence encodes Human-pro-Asp 2(a) delta TM
 CC (low GC), a synthetic version of Asp 2(a) whose GC content has been
 CC altered to facilitate expression in E.coli

SO Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 2.67e-06 Length: 1302
 Score: 16.00 Matches: 16
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x AAS11528 (1-1302)

Qy 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyVala 16
 |||
 4 ACTGACATGATTCGTCTGCGCACTGCGTACGCGTCTGGTGTCT 51

RESULT 15
 ABL52468 ID ABL52468 standard; cDNA; 1302 BP.
 XX
 AC ABL52468;
 XX
 DT 16-JUL-2002 (first entry)
 XX
 DE Human-pro-Asp-2(a)deltaTM (low GC) nucleotide sequence SEQ ID NO:25.
 XX
 KW Human: Asp-1; Asp-2; aspartyl protease; Alzheimer's disease; proteolytic;
 KW amyloid precursor protein; APP; gene; ss.
 XX
 OS Homo sapiens.
 OS
 FH Key Location/Qualifiers
 FT CDS 1..1302
 FT /tag= a
 FT /product= "human-pro-Asp-2(a)deltaTM (low GC)"

GB2367060-A.
 XX
 PN 27-MAR-2002.
 XX
 PD 29-OCT-2001; 2001GB-00025934.
 XX
 PF 23-SEP-1999; 99US-00404133.
 PR 23-SEP-1999; 99US-0155493P.
 PR 23-SEP-1999; 99WO-US020881.
 PR 13-OCT-1999; 99US-00416901.
 PR 06-DEC-1999; 99US-0169232P.
 PR 22-SEP-2000; 2000GB-00023315.
 XX
 PA (PHAA) PHARMACIA & UPJOHN CO.
 PA
 PI Bienkowski MJ, Gurney M;
 PI
 DR MPI; 2002-397167/43.
 DR P-PSDB; ABB78601.
 XX
 PT Human aspartyl protease 1 substrates useful in assays to detect aspartyl
 PT protease activity, e.g. for the diagnosis of Alzheimer's disease.
 XX

PS Example 9; Fig 8; 182pp; English.

XX
CC The present invention describes a human aspartyl protease 1 (hu-Asp1)
CC substrate (I) which comprises a peptide of no more than 50 amino acids,
CC and which comprises the 8 amino acid sequence Gly-Leu-Ala-Leu-Ala-Leu-
CC Glu-Pro. Also described are: (1) a method (II) for assaying hu-Asp1
CC proteolytic activity, comprising: (a) contacting a hu-Asp1 protein with
CC (1) under acidic conditions; and (b) determining the level of hu-Asp1
CC proteolytic activity; (2) a purified polynucleotide (III) comprising a
CC nucleotide sequence that hybridises under stringent conditions to the non
CC -coding strand complementary to a defined 1804 nucleotide sequence (see
CC ABU52456) where the nucleotide sequence encodes a polypeptide having Asp1
CC proteolytic activity and lacks nucleotides encoding a transmembrane
CC domain; (3) a purified polynucleotide (III') comprising a sequence that
CC hybridises under stringent conditions to (III) (the nucleotide sequence
CC encodes a polypeptide further lacking a pro-peptide domain corresponding
CC to amino acids 23-62 of hu-Asp1 (see AB078589)); (4) a vector (IV)
CC comprising (III) or (III'); and (5) a host cell (V) transformed or
CC transfected with (III), (III') and/or (IV). The hu-Asp1 protease
CC substrate (1) may be used as an enzyme substrate in assays to detect
CC aspartyl protease activity, (II) and therefore diagnose diseases
CC associated with aberrant hu-Asp1 expression and activity such as
CC Alzheimer's disease. Hu-Asp1 has been localised to chromosome 21, while
CC hu-Asp2 has been localised to chromosome 11q23.3-24.1. The present
CC sequence encodes human-pro-Asp-2(a)delatam (low GC), which is given in an
CC example from the present invention
XX

SQ Sequence 1302 BP; 281 A; 367 C; 370 G; 284 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	2	67e-06	Length:	1302
Score:	16.00		Matches:	16
Percent Similarity:	100.00%		Conservative:	0
Best Local Similarity:	100.00%		Mismatches:	0
Query Match:	100.00%		Indels:	0
DB:	6		Gaps:	0

US-10-726-967A-3 (1-16) x ABU52468 (1-1302)

QY	1	ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyAla	16
DB	4	ACTCAGCATGGTATTGCTCGCACTGCGTAGCGGTCTGGGTGGTCT	51

Search completed: July 27, 2005, 19:10:44
Job time : 434 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 18:52:49 ; Search time 134 Seconds

(without alignments)
195.376 Million cell updates/sec

Title: US-10-726-967A-3

Perfect score: 16
Sequence: 1 TQHGRLPLRSLGCGA 16

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Ygapop 60.0 ,	Ygapext 60.0
Fgapop 6.0 ,	Fgapext 7.0
Delop 6.0 ,	Delext 7.0

Searched: 1202784 seqs, 818138359 residues

Word size: 1

Total number of hits satisfying chosen parameters: 2397162

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

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-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi
-LIST=45 -DOCLISTN=200 -THR SCORE=quality -THR MIN=1 -ALIGN=15 -MODE=LOCAL
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Database :

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	1287	3	US-09-548-372D-50
2	16	100.0	1287	3	US-09-548-367D-50
3	16	100.0	1287	3	US-09-548-367D-50
4	16	100.0	1287	4	US-09-551-853D-50
5	16	100.0	1287	4	US-09-548-376D-50
6	16	100.0	1287	4	US-09-548-376D-50
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8	16	100.0	1287	4	US-09-548-373D-50
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11	16	100.0	1287	4	US-09-548-368D-50
12	16	100.0	1287	4	US-09-794-925A-50

13	16	100.0	1302	3	US-09-548-372D-25	Sequence 25, Appl
14	16	100.0	1302	3	US-09-548-367D-25	Sequence 25, Appl
15	16	100.0	1302	3	US-09-551-853D-25	Sequence 25, Appl
16	16	100.0	1302	4	US-09-416-901B-25	Sequence 25, Appl
17	16	100.0	1302	4	US-09-548-376D-25	Sequence 25, Appl
18	16	100.0	1302	4	US-09-794-927A-25	Sequence 25, Appl
19	16	100.0	1302	4	US-09-548-373D-25	Sequence 25, Appl
20	16	100.0	1302	4	US-09-795-847B-25	Sequence 25, Appl
21	16	100.0	1302	4	US-09-869-414-25	Sequence 25, Appl
22	16	100.0	1302	4	US-09-548-366F-25	Sequence 25, Appl
23	16	100.0	1302	4	US-09-548-368D-25	Sequence 25, Appl
24	16	100.0	1302	4	US-09-794-925A-25	Sequence 25, Appl
25	16	100.0	1302	4	US-09-806-194A-25	Sequence 25, Appl
26	16	100.0	1305	3	US-09-548-372D-52	Sequence 52, Appl
27	16	100.0	1305	3	US-09-548-367D-52	Sequence 52, Appl
28	16	100.0	1305	4	US-09-551-853D-52	Sequence 52, Appl
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30	16	100.0	1305	4	US-09-548-376D-52	Sequence 52, Appl
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32	16	100.0	1305	4	US-09-548-373D-52	Sequence 52, Appl
33	16	100.0	1305	4	US-09-795-847B-52	Sequence 52, Appl
34	16	100.0	1305	4	US-09-869-414-52	Sequence 52, Appl
35	16	100.0	1305	4	US-09-548-366F-52	Sequence 52, Appl
36	16	100.0	1305	4	US-09-548-368D-52	Sequence 52, Appl
37	16	100.0	1305	4	US-09-794-925A-52	Sequence 52, Appl
38	16	100.0	1341	3	US-09-548-372D-21	Sequence 21, Appl
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43	16	100.0	1341	4	US-09-794-927A-21	Sequence 21, Appl
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45	16	100.0	1341	4	US-09-795-847B-21	Sequence 21, Appl

ALIGNMENTS

RESULT 1
US-09-548-372D-50
; Sequence 50, Application US/09548372D
; Patent No. 6420534
; GENERAL INFORMATION:
; APPLICANT: GUNNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 29915/62801
; CURRENT APPLICATION NUMBER: US/09/548,372D
; CURRENT FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-372D-50
Alignment Scores:
Pred. No.: 1.48e-06
Score: 16.00
Percent Similarity: 100.00%
Best Local Similarity: 100.00%
Query Match: 100.00%
DB: 3
Length: 1287
Matches: 16
Conservative: 0
Mismatch: 0
Indels: 0
Gaps: 0

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QY 1 ThrglnhsglyllleargleuProleuArgserglyleuglygla 16
Db 64 ACCGAGCAGCGCATCCGGCTGCCCTCGCGAGCGGCGCTGGGGGGCGCC 111

RESULT 2
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; Sequence 50, Application US/09548367D
; Patent No. 6440698
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280H
; CURRENT APPLICATION NUMBER: US/09/548,367D
; PRIOR FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-548-367D-50

Alignment Scores:
Pred. No.: 1,48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-548-367D-50 (1-1287)
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Db 64 ACCGAGCAGCGCATCCGGCTGCCCTCGCGAGCGGCGCTGGGGGGCGCC 111

RESULT 3
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; Patent No. 6500667
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280L
; CURRENT APPLICATION NUMBER: US/09/551,853D
; PRIOR FILING DATE: 2000-04-18
; PRIOR APPLICATION NUMBER: US 60/155,493
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; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
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; TYPE: DNA
; ORGANISM: Artificial sequence
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; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-551-853D-50

Alignment Scores:
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Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-416-901B-50 (1-1287)
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Db 64 ACCGAGCAGCGCATCCGGCTGCCCTCGCGAGCGGCGCTGGGGGGCGCC 111

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; Patent No. 6699671
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280A
; CURRENT APPLICATION NUMBER: US/09/416,901B
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
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; PRIOR FILING DATE: 1999-09-23
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; PRIOR FILING DATE: 1998-09-24
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; ORGANISM: Artificial sequence
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; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-416-901B-50

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Pred. No.: 1,48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
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US-10-726-967A-3 (1-16) x US-09-416-901B-50 (1-1287)
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Db 64 ACCGAGCAGCGCATCCGGCTGCCCTCGCGAGCGGCGCTGGGGGGCGCC 111

RESULT 5
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; Patent No. 6706485
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280F
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CURRENT APPLICATION NUMBER: US/09/548,376D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
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PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 50
LENGTH: 1287
TYPE: DNA
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-376D-50

Alignment Scores:
Pred. No.: 1,48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
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US-10-726-967A-3 (1-16) x US-09-548-376D-50 (1-1287)
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Db 64 ACCGACGCGGATCCGGCTGCCCTGCGAGCGGCTGGGGGGCGCC 111

RESULT 6
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Patent No. 6727074
GENERAL INFORMATION:
APPLICANT: Gurney et al.
TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
FILE REFERENCE: 29915/6280FG
CURRENT APPLICATION NUMBER: US/09/794,927A
CURRENT FILING DATE: 2001-02-27
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
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TYPE: DNA
ORGANISM: Artificial sequence
FEATURE:
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US-09-794-927A-50

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Pred. No.: 1,48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-794-927A-50 (1-1287)
QY 1 ThrglnHieGlylleArgLeuProLeuArgSerGlyLeuGlyAla 16
Db 64 ACCGACGCGGATCCGGCTGCCCTGCGAGCGGCTGGGGGGCGCC 111

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US-09-548-373D-50
Sequence 50, Application US/09548373D
Patent No. 6737510
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
FILE REFERENCE: 29915/6280B
CURRENT APPLICATION NUMBER: US/09/548,373D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 50
LENGTH: 1287
TYPE: DNA
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2(b) delta TM
US-09-548-373D-50

Alignment Scores:
Pred. No.: 1,48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-548-373D-50 (1-1287)
QY 1 ThrglnHieGlylleArgLeuProLeuArgSerGlyLeuGlyAla 16
Db 64 ACCGACGCGGATCCGGCTGCCCTGCGAGCGGCTGGGGGGCGCC 111

RESULT 8
US-09-795-847B-50
Sequence 50, Application US/09795847B
Patent No. 6753163
GENERAL INFORMATION:
APPLICANT: Gurney, Mark E.
APPLICANT: Bienkowski, Michael J.
APPLICANT: Heinrikson, Robert L.
APPLICANT: Parodi, Luis A.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
FILE REFERENCE: 28341/6280DE
CURRENT APPLICATION NUMBER: US/09/795,847B
CURRENT FILING DATE: 2001-02-28
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24

PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-795-847B-50

Alignment Scores:
Pred. No.: 1.48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-795-847B-50 (1-1287)

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RESULT 9
US-09-869-414-50
; Sequence 50, Application US/09869414
; Patent No. 6790610
; GENERAL INFORMATION:
; APPLICANT: Beinkowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M
; CURRENT APPLICATION NUMBER: US/09/869,414
; PRIOR FILING DATE: 2001-06-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; OTHER INFORMATION: delta TM
US-09-869-414-50

Alignment Scores:
Pred. No.: 1.48e-06 Length: 1287
Score: 16.00 Matches: 16
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Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-869-414-50 (1-1287)

Qy 1 ThrglnhsglyileargleuProleuArgserglyleuglyyala 16
Db 64 ACCGACGCGGCAATCCGGCTGCCCTGCGACGCGGCTGGGGGGCGCC 111

RESULT 10
US-09-548-366F-50

; Sequence 50, Application US/09548366F
; Patent No. 6797487
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280J
; CURRENT APPLICATION NUMBER: US/09/548,366F
; PRIOR FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-548-366F-50

Alignment Scores:
Pred. No.: 1.48e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-548-366F-50 (1-1287)

Qy 1 ThrglnhsglyileargleuProleuArgserglyleuglyyala 16
Db 64 ACCGACGCGGCAATCCGGCTGCCCTGCGACGCGGCTGGGGGGCGCC 111

RESULT 11
US-09-548-368D-50
; Sequence 50, Application US/09548368D
; Patent No. 6825023
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280C
; CURRENT APPLICATION NUMBER: US/09/548,368D
; PRIOR FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-548-368D-50

Alignment Scores:
Pred. No.: 1.48e-06 Length: 1287
Score: 16.00 Matches: 16

Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
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US-10-726-967A-3 (1-16) x US-09-548-368D-50 (1-1287)

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DB 64 ACCCAGCAGCGCATCCGGCTGCCCTCGCCAGCGGCTGGGGGGCCGCC 111

RESULT 12

US-09-794-925A-50
Sequence 50, Application US/09794925A

Patent No. 6828117

GENERAL INFORMATION:

APPLICANT: Gurney et al.

TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses

FILE REFERENCE: 29915/6280H1

CURRENT APPLICATION NUMBER: US/09/794,925A

CURRENT FILING DATE: 2001-02-27

PRIOR FILING DATE: 1999-10-13

PRIOR APPLICATION NUMBER: 60/155,493

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: 09/404,133

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: PCT/US99/20881

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: 60/101,594

PRIOR FILING DATE: 1998-09-24

NUMBER OF SEQ ID NOS: 74

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 50

LENGTH: 1287

TYPE: DNA

ORGANISM: Artificial sequence

FEATURE: OTHER INFORMATION: Hu-Asp2 (b) delta TM

US-09-794-925A-50

Alignment Scores:

Pred. No.: 1,48e-06 Length: 1287

Score: 16.00 Matches: 16

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 100.00% Indels: 0

DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-794-925A-50 (1-1287)

QY 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyGlyVala 16
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DB 64 ACCCAGCAGCGCATCCGGCTGCCCTCGCCAGCGGCTGGGGGGCCGCC 111

RESULT 13

US-09-548-372D-25
Sequence 25, Application US/09548372D

Patent No. 6420534

GENERAL INFORMATION:

APPLICANT: Gurney et al.

TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES

FILE REFERENCE: 29915/6280I

CURRENT APPLICATION NUMBER: US/09/548,372D

CURRENT FILING DATE: 2000-04-12

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: US 09/404,133

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: PCT/US99/20881

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 25
LENGTH: 1302
TYPE: DNA
ORGANISM: Homo sapiens
US-09-548-372D-25

Alignment Scores:

Pred. No.: 1.5e-06 Length: 1302

Score: 16.00 Matches: 16

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 100.00% Indels: 0

DB: 3 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-548-372D-25 (1-1302)

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RESULT 14

US-09-548-367D-25
Sequence 25, Application US/09548367D

Patent No. 6440698

GENERAL INFORMATION:

APPLICANT: Gurney et al.

TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES

FILE REFERENCE: 29915/6280H

CURRENT APPLICATION NUMBER: US/09/548,367D

CURRENT FILING DATE: 2000-04-12

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: US 09/404,133

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: PCT/US99/20881

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: US 60/101,594

PRIOR FILING DATE: 1998-09-24

NUMBER OF SEQ ID NOS: 73

SOFTWARE: PatentIn version 3.1

SEQ ID NO 25

LENGTH: 1302

TYPE: DNA

ORGANISM: Homo sapiens

US-09-548-367D-25

Alignment Scores:

Pred. No.: 1.5e-06 Length: 1302

Score: 16.00 Matches: 16

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 100.00% Indels: 0

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US-10-726-967A-3 (1-16) x US-09-548-367D-25 (1-1302)

QY 1 ThrGlnHisGlyIleArgLeuProLeuArgSerGlyLeuGlyGlyVala 16
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DB 4 ACTCAGCATGTATTGCTGCTGCCACTGCGTACCGGCTGGGTGCT 51

RESULT 15

US-09-551-853D-25
Sequence 25, Application US/09551853D

Patent No. 650867

GENERAL INFORMATION:

APPLICANT: Gurney et al.

TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES

FILE REFERENCE: THEREOF

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FILE REFERENCE: 29915/6280L
CURRENT APPLICATION NUMBER: US/09/551,853D
PRIOR FILING DATE: 2000-04-18
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 25
LENGTH: 1302
TYPE: DNA
ORGANISM: Homo sapiens
US-09-551-853D-25

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OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 19:03:29 ; Search time 625 Seconds

(without alignments)
165.537 Million cell updates/sec

Title: US-10-726-967A-3

Perfect score: 16

Sequence: 1 TQHGRLPLRSLGCA 16

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Published Applications NA:
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and is derived by analysis of the total score distribution.

SUMMARIES

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4	16	100.0	1287	9	US-09-794-748-50	Sequence 50, Appl
5	16	100.0	1287	9	US-09-794-925-50	Sequence 50, Appl
6	16	100.0	1287	9	US-09-681-442-50	Sequence 50, Appl
7	16	100.0	1287	10	US-09-869-414-50	Sequence 50, Appl
8	16	100.0	1287	10	US-09-548-366-50	Sequence 50, Appl
9	16	100.0	1287	18	US-10-652-927-50	Sequence 50, Appl
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38	16	100.0	1305	19	US-10-652-045-52	Sequence 52, Appl
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44	16	100.0	1341	9	US-09-794-748-21	Sequence 21, Appl
45	16	100.0	1341	9	US-09-794-925-21	Sequence 21, Appl

ALIGNMENTS

RESULT 1
US-09-794-927-50
; Sequence 50, Application US/09794927
; Patent No. US20010016324A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Blenkow, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Ridiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: US95
; FILE REFERENCE: 28341/6280FG
; CURRENT APPLICATION NUMBER: US/09/794, 927
; CURRENT FILING DATE: 2001-02-27

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; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; US-09-794-927-50

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Alignment Scores:

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Score:	16.00	Matches:	16
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
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US-10-726-967a-3 (1-16) x US-09-794-927-50 (1-1287)

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Db      64 ACCGACGACGCGATCCGCGCTCCCTCGCGACGCGGCTCGGGGGGCC 111

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RESULT 2

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; Patent No. US20010018208A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; FILE REFERENCE: 28341/6280DE
; CURRENT APPLICATION NUMBER: US/09/795,847
; CURRENT FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; US-09-795-847-50

```

```

; Alignment Scores:
; US-09-795-847-50
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; OTHER INFORMATION: delta TM
; US-09-795-847-50

```

Pred. No.:	2.94e-06	Length:	1287
Score:	16.00	Matches:	16
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	9	Gaps:	0

US-10-726-967a-3 (1-16) x US-09-795-847-50 (1-1287)

```

QY      1 ThrglnHsglyYlleArgLeuProLeuAArgSerGlyLeuGlyGlyAla 16
Db      64 ACCGACGACGCGATCCGCGCTCCCTCGCGACGCGGCTCGGGGGGCC 111

```

RESULT 3

```

; Sequence 50, Application US/09794743
; Patent No. US20010021391A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; FILE REFERENCE: 28341/6280BC
; CURRENT APPLICATION NUMBER: US/09/794,743
; CURRENT FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; US-09-794-743-50

```

Alignment Scores:

Pred. No.:	2.94e-06	Length:	1287
Score:	16.00	Matches:	16
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	9	Gaps:	0

US-10-726-967a-3 (1-16) x US-09-794-743-50 (1-1287)

```

QY      1 ThrglnHsglyYlleArgLeuProLeuAArgSerGlyLeuGlyGlyAla 16
Db      64 ACCGACGACGCGATCCGCGCTCCCTCGCGACGCGGCTCGGGGGGCC 111

```

RESULT 4

```

; Sequence 50, Application US/09794748
; Patent No. US20020037315A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.

```

```

; Alignment Scores:
; US-09-794-748-50
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; OTHER INFORMATION: delta TM
; US-09-794-748-50

```

```

; APPLICANT: Yan, Rigdang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280JL
; CURRENT APPLICATION NUMBER: US/09/794,748
; CURRENT FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Ap2 (b)
; US-09-794-748-50

Alignment Scores:
Pred. No.: 2,94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-10-726-967a-3 (1-16) x US-09-794-748-50 (1-1287)

Qy 1 ThrglnHieglYlIeargleuPfoleuArgserGlyLeuGlYglYAla 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTGCGCAGCGGCGCTGGGGGCGGCC 111

RESULT 5
US-09-794-925-50
; Sequence 50, Application US/09794925
; Patent No. US20020064819A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Rigdang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280HI
; CURRENT APPLICATION NUMBER: US/09/794,925
; CURRENT FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
```

```

; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Ap2 (b)
; OTHER INFORMATION: delta TM
; US-09-794-925-50

Alignment Scores:
Pred. No.: 2,94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-10-726-967a-3 (1-16) x US-09-794-925-50 (1-1287)

Qy 1 ThrglnHieglYlIeargleuPfoleuArgserGlyLeuGlYglYAla 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTGCGCAGCGGCGCTGGGGGCGGCC 111

RESULT 6
US-09-681-442-50
; Sequence 50, Application US/09681442
; Patent No. US20020081634A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Rigdang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280FG
; CURRENT APPLICATION NUMBER: US/09/681,442
; CURRENT FILING DATE: 2001-04-05
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Ap2 (b)
; OTHER INFORMATION: delta TM
; US-09-681-442-50

Alignment Scores:
Pred. No.: 2,94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-10-726-967a-3 (1-16) x US-09-681-442-50 (1-1287)

Qy 1 ThrglnHieglYlIeargleuPfoleuArgserGlyLeuGlYglYAla 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTGCGCAGCGGCGCTGGGGGCGGCC 111

RESULT 7
US-09-869-414-50
; Sequence 50, Application US/09869414
; Publication No. US20030077226A1
```

```

; GENERAL INFORMATION:
; APPLICANT: Bienkowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M
; CURRENT APPLICATION NUMBER: US/09/869,414
; PRIOR FILING DATE: 2001-06-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
US-09-869-414-50

Alignment Scores:
Pred. No.: 2,94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 10 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-869-414-50 (1-1287)
QY 1 ThrglnhlglylleaRgleuProleuArgSerGlyLeuGlyGlyala 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTCGCGCAGCGGCTGGGGGGCGCC 111

RESULT 8
US-09-548-366-50
; Sequence 50, Application US/09548366
; Publication No. US20030104365A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; FILE REFERENCE: 28341/6280A
; CURRENT APPLICATION NUMBER: US/09/548,366
; PRIOR FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 65
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2(b)
```

```

; OTHER INFORMATION: delta TM
US-09-548-366-50

Alignment Scores:
Pred. No.: 2,94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 10 Gaps: 0

US-10-726-967A-3 (1-16) x US-09-548-366-50 (1-1287)
QY 1 ThrglnhlglylleaRgleuProleuArgSerGlyLeuGlyGlyala 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTCGCGCAGCGGCTGGGGGGCGCC 111

RESULT 9
US-10-652-927-50
; Sequence 50, Application US/10652927
; Publication No. US20040043408A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 28915/6280N3
; CURRENT APPLICATION NUMBER: US/10/652,927
; PRIOR FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: 09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2(b) delta TM
US-10-652-927-50

Alignment Scores:
Pred. No.: 2,94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 18 Gaps: 0

US-10-726-967A-3 (1-16) x US-10-652-927-50 (1-1287)
QY 1 ThrglnhlglylleaRgleuProleuArgSerGlyLeuGlyGlyala 16
Db 64 ACCCAGCAGCGCATCCGGCTGCCCTCGCGCAGCGGCTGGGGGGCGCC 111

RESULT 10
US-10-652-830-50
; Sequence 50, Application US/10652830
; Publication No. US20040048303A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280N1
```

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; CURRENT APPLICATION NUMBER: US/10/652,830
; CURRENT FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: 09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-10-652-830-50

Alignment Scores:
Pred. No.:      2,94e-06      Length:      1287
Score:          16.00         Matches:      16
Percent Similarity: 100.00%   Conservaive:  0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    100.00%      Indels:      0
DB:             18           Gaps:       0

US-10-726-967a-3 (1-16) x US-10-652-830-50 (1-1287)

QY      1  ThrGlnHlaGlyIleArgLeuProLeuArgSerGlyLeuGlyGlyAla 16
Db      64  ACCCAGCAGCGCATCCGCTGCCCTCGCCAGCGGCTCGGGGGGCCCC 111

RESULT 11
US-10-652-045-50
; Sequence 50, Application US/10652045
; Publication No. US20040166507A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280N2
; CURRENT APPLICATION NUMBER: US/10/652,045
; CURRENT FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: 09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-10-652-045-50

Alignment Scores:
Pred. No.:      2,94e-06      Length:      1287
Score:          16.00         Matches:      16
Percent Similarity: 100.00%   Conservaive:  0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    100.00%      Indels:      0
DB:             18           Gaps:       0
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```

Score:          16.00         Matches:      16
Percent Similarity: 100.00%   Conservaive:  0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    100.00%      Indels:      0
DB:             19           Gaps:       0

US-10-726-967a-3 (1-16) x US-10-652-045-50 (1-1287)

QY      1  ThrGlnHlaGlyIleArgLeuProLeuArgSerGlyLeuGlyGlyAla 16
Db      64  ACCCAGCAGCGCATCCGCTGCCCTCGCCAGCGGCTCGGGGGGCCCC 111

RESULT 12
US-10-476-935-50
; Sequence 50, Application US/10476935
; Publication No. US20040234976A1
; GENERAL INFORMATION:
; APPLICANT: Beinkowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M1
; CURRENT APPLICATION NUMBER: US/10/476,935
; CURRENT FILING DATE: 2003-11-06
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
US-10-476-935-50

Alignment Scores:
Pred. No.:      2,94e-06      Length:      1287
Score:          16.00         Matches:      16
Percent Similarity: 100.00%   Conservaive:  0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    100.00%      Indels:      0
DB:             20           Gaps:       0

US-10-726-967a-3 (1-16) x US-10-476-935-50 (1-1287)

QY      1  ThrGlnHlaGlyIleArgLeuProLeuArgSerGlyLeuGlyGlyAla 16
Db      64  ACCCAGCAGCGCATCCGCTGCCCTCGCCAGCGGCTCGGGGGGCCCC 111

RESULT 13
US-10-477-076-50
; Sequence 50, Application US/10477076
; Publication No. US20050080232A1
; GENERAL INFORMATION:
; APPLICANT: Beinkowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M2
; CURRENT APPLICATION NUMBER: US/10/477,076
; CURRENT FILING DATE: 2003-11-06
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
```

```

; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Asp2 (b)
; OTHER INFORMATION: delta TM
US-10-477-076-50

Alignment Scores:
Pred. No.: 2.94e-06 Length: 1287
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 21 Gaps: 0

US-10-726-967a-3 (1-16) x US-10-477-076-50 (1-1287)

Qy 1 Thrglnhsglylleargleuproleuargserglyleuglyala 16
Db 64 ACCGACGCGGCGATCGGCTGCCCTGGCGAGCGGCGCTGGGGGCGGCC 111

RESULT 14
US-09-794-927-25
; Sequence 25, Application US/09794927
; Patent No. US20010016324A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Helmricks, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; TITLE OF INVENTION: THEREFOR
; FILE REFERENCE: 28341/6280FG
; CURRENT APPLICATION NUMBER: US/09/794,927
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 25
; LENGTH: 1302
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-794-927-25

Alignment Scores:
Pred. No.: 2.97e-06 Length: 1302
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967a-3 (1-16) x US-09-795-847-25 (1-1302)

Qy 1 Thrglnhsglylleargleuproleuargserglyleuglyala 16
Db 4 ACTGACGATGGTATTCGTCTGCCACTGCGTAGCGGCTCGGGTGCT 51

Search completed: July 27, 2005, 20:48:51
Job time : 626 secs
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GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 18:50:24 ; Search time 3143 Seconds
(without alignments)
193.773 Million cell updates/sec

Title: US-10-726-967A-3

Perfect score: 16

Sequence: 1 TGHGRLPLRSLGCA 16

Scoring table:

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Ygapop 60.0 , Ygapext 60.0
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 34239544 seqs, 19032134700 residues

Word size: 1

Total number of hits satisfying chosen parameters: 68473592

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Command line parameters:

-MODE=frame+pn.model -DEV=xlp
-Q/cn2.1/USPTO.spool.p/US10726967/runat.26072005.130816.6487/app.query.fasta.1.199
-DB=EST -OFMT=fasta -SUFFIX=olip2n.rst -MINMATCH=0.1 -LOOPL=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi -LIST=45
-DOCALLIGN=200 -THR.SCORE=quality -THR.MIN=1 -ALIGN=15 -MODE=LOCAL -OUTFMT=prc
-NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US10726967 -@CN 1 1 5180 -@runat.26072005.130816.6487 -NCPU=6 -ICPU=3
-NO MMAP -LARGEOUTERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-BEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

Database :

EST.*
1: gb_est1.*
2: gb_est2.*
3: gb_hic.*
4: gb_est3.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_gse1.*
9: gb_gse2.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	611	7	CN484125 hw42d08.y
2	16	100.0	1506	9	AY417360 Homo sapi
3	13	81.2	424	4	B1337739 361259 MA
4	13	81.2	563	4	BG833894 351953 MA
5	13	81.2	616	5	BP452814 BP452814
6	11	68.8	171	9	CE200035 C19T-g8s-
7	10	62.5	493	7	CN692524 E0324B10-
8	10	62.5	533	7	CN697484 E0394H10-
9	10	62.5	547	7	CP903755 A0413D10-

10	10	62.5	563	7	CP906581
11	10	62.5	576	7	CF171218
12	10	62.5	600	6	BY713879
13	10	62.5	619	2	BB644736
14	10	62.5	637	6	CD348605
15	10	62.5	639	2	BB632244
16	10	62.5	639	2	BB652612
17	10	62.5	640	6	BY724111
18	10	62.5	650	2	BB624169
19	10	62.5	653	6	BY727440
20	10	62.5	673	2	BB640442
21	10	62.5	673	2	BB650860
22	10	62.5	727	6	CA749486
23	10	62.5	836	5	BK374914
24	10	62.5	844	1	AL544727
25	10	62.5	1123	5	BX376891
26	10	62.5	1506	9	AY117362
27	10	62.5	3634	3	AK041285
28	10	62.5	3859	3	AK014464
29	10	62.5	3877	3	AK033112
30	10	62.5	4046	3	AK049626
31	10	62.5	4048	3	AK082317
32	10	62.5	4101	3	AK046175
33	10	62.5	409	6	CB215573
34	9	56.2	566	8	BH881298
35	9	56.2	913	9	CG324082
36	9	56.2	986	5	BX393934
37	9	56.2	1765	2	BF572300
38	9	56.2	156	7	CO060378
39	8	50.0	220	7	CF681698
40	8	50.0	234	2	BB569323
41	8	50.0	258	2	BF872950
42	8	50.0	272	4	BM258402
43	8	50.0	295	4	BM574145
44	8	50.0	305	9	CC810341
45	8	50.0			

ALIGNMENTS

RESULT 1
CN484125 611 bp mRNA linear EST 26-APR-2004
hw42d08.y2 Human primary human ocular pericytes. Unamplified (hw)
DEFINITION Homo sapiens CDNA clone hw42d08 5', mRNA sequence.
LOCUS CN484125
ACCESSION CN484125 GI:46565629
VERSION
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
AUTHORS Tsai, J.Y. and Wistow, G.
TITLE 1 (bases 1 to 611)
EXPRESSION TAG ANALYSIS OF CULTURED PRIMARY HUMAN OCULAR PERICYTES

JOURNAL Unpublished (2004)
COMMENT Contact: Wistow G
Section on Molecular Structure and Function
National Eye Institute
6/331, NIH, Bethesda, MD 20892-2740, USA
Tel: 301 402 3452
Fax: 301 496 0078
Email: gwaem@nei.nih.gov
Plate: 42 row: d column: 08
Seg primer: M13Rpl reverse primer (ABI).
Location/Qualifiers

FEATURES
source 1..611
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="hw42d08"
/cell_type="pericytes"

/dev_stage="Adult"
/lab_host="EMDH10B"
/clone_lib="Human primary human ocular pericytes.
Unamplified (hw)"
/note="Organ: Eye; Vector: pSPORT1; RNA was extracted from
primary human pericytes in culture. A directionally cloned
cDNA library in the pSPORT vector (Invitrogen) was
constructed at Bioserve Biotechnology (Laurel MD)
essentially following the protocols of the Superscript
Plasmid System full details of which are contained in the
manufacturer's instruction manual
(http://www.lifetechn.com/). First strand synthesis was
carried out using a Not I primer-adaptor
[5'-pGACTAGTCTAGATCGAGCGCGCC(T)15-3']. cDNA was
cloned in Not I/Sal I sites. EST analysis was performed at
the NIH Intramural Sequencing Center (NISC)."

ORIGIN

Alignment Scores:

Pred. No.: 9,47e-06 Length: 611
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 7 Gaps: 0

US-10-726-967A-3 (1-16) x CM484125 (1-611)

QY 1 ThrGlnH1eG1Y1leArgLeuProleuArgSerG1yleuG1yG1yAla 16

Db 381 ACCCGACGCGCATCGCGCTGCCCTGCCAGCGGCGCTGGGGGCGCC 428

RESULT 2

AY417360

LOCUS AY417360 1506 bp DNA linear GSS 17-DEC-2003
DEFINITION Homo sapiens BACE gene, VIRUTAL TRANSCRIPT, partial sequence.
ACCESSION AY417360
VERSION AY417360.1 GI:39773320
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 1506)
Clark,A.G., Gnanowski,S., Nielson,R., Thomas,P., Kejarival,A.,
Todd,M.A., Tanenbaum,D.M., Civello,D.R., Lu,F., Murphy,B.,
Fertiera,S., Wang,G., Zheng,X.H., White,T.J., Sninsky,J.J.,
Adams,M.D. and Cargill,M.
Inferring nonneutral evolution from human-chimp-mouse orthologous
gene trios

AUTHORS

TITLE

JOURNAL Science 302 (5652), 1960-1963 (2003)
PUBMED 14671302
2 (bases 1 to 1506)

REFERENCE

AY417360

AUTHORS

Clark,A.G., Gnanowski,S., Nielson,R., Thomas,P., Kejarival,A.,
Todd,M.A., Tanenbaum,D.M., Civello,D.R., Lu,F., Murphy,B.,
Fertiera,S., Wang,G., Zheng,X.H., White,T.J., Sninsky,J.J.,
Adams,M.D. and Cargill,M.
Direct Submission
Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
Rockville, MD 20850, USA

COMMENT

This sequence was made by sequencing genomic exons and ordering
them based on alignment.

FEATURES

source

gene

ORIGIN

/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/gene="BACE"
/locus_tag="HCM6198"

Alignment Scores:

Pred. No.: 2,11e-05 Length: 1506
Score: 16.00 Matches: 16
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967A-3 (1-16) x AY417360 (1-1506)

QY 1 ThrGlnH1eG1Y1leArgLeuProleuArgSerG1yleuG1yG1yAla 16

Db 64 ACCCGACGCGCATCGCGCTGCCCTGCCAGCGGCGCTGGGGGCGCC 111

RESULT 3

B1337739

LOCUS B1337739 424 bp mRNA linear EST 30-JUL-2001
DEFINITION 361259 MARC 1P1G Sus scrofa cDNA 5', mRNA sequence.
ACCESSION B1337739
VERSION B1337739.1 GI:15031022
KEYWORDS EST.
SOURCE Sus scrofa (pig)
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Sus.

REFERENCE 1 (bases 1 to 424)
Fahrenkrug,S.C., Smith,T.P.L., Freking,B.A., Cho,J., White,J.,
Vallet,J., Wise,T., Rohrer,G.A., Pertea,G., Sultana,R.,
Quackenbush,J. and Keefe,J.W.
Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly

TITLE

JOURNAL Mamm. Genome 13 (6), 475-478 (2002)
MEDLINE 22213789
PUBMED 12226715

COMMENT

Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@mail.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -mismatch 18
and -mismatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCACGACGACGACG
Plate: 126 row: K column: 16
Seq primer: ATTAGGTGACACTATAG.
Location/Qualifiers
1..424

FEATURES

source

/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 1P1G"
/note="Vector: pCMV SPORT6; Site 1: NotI, Site 2: SalI;
library made from pooled tissue from day 11, 13, 15, 20,
and 30 embryos."

ORIGIN

Alignment Scores:

Pred. No.: 0.00572 Length: 424
Score: 13.00 Matches: 13
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 81.25% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) x B1337739 (1-424)

QY

4 GlyIleArgLeuProleuArgSerG1yleuG1yG1yAla 16

|||||

DB 376 GGCATCCGGCTGCCCTGCGAAGCGGCTTGGGGGGCA 414

RESULT 4
LOCUS BG833894 563 bp mRNA linear EST 25-MAY-2001
DEFINITION BG833894 351953 MARC 1P1G Sus scrofa CDNA 5', mRNA sequence.
ACCESSION BG833894
VERSION BG833894.1 GI:14198715
KEYWORDS EST.
SOURCE Sus scrofa (pig)
ORGANISM Sus scrofa
REFERENCE Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Eukaryota; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
1 (bases 1 to 563)
Fahnenkrug, S.C., Smith, T.P.L., Freking, B.A., Cho, J., White, J., Valler, J., Wise, T., Rohrer, G.A., Ferreira, G., Sultana, R., Quackenbush, J. and Keele, J.W.
Porcine gene discovery by normalized cDNA-library sequencing and EST cluster assembly
Mamm. Genome 13 (8), 475-478 (2002)

TITLE Mamm. Genome 13 (8), 475-478 (2002)

JOURNAL MEDLINE 22213789
PUBMED 12226715
COMMENT Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred v0.980904.e. Vector identified by cross_match with the -minscore 18 and -minmatch 12 options.
PCR Primers
FORWARD: AGGAACACAGCTATGACCAT
BACKWARD: GTTTCACGATCAGCAGC
Place: 111 row: G column: 20
Seq primer: ATTAGTGACACTATAG.
Location/Qualifiers
1..563
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/cruise_type="pooled"
/lab_host="DH10B"
/clone_11b="MARC 1P1G"
/note="Vector: PCMV SPORT6; Site 1: NotI; Site 2: SalI; library made from pooled tissue from day 11, 13, 15, 20, and 30 embryos."

ORIGIN
source
1..563
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/cruise_type="pooled"
/lab_host="DH10B"
/clone_11b="MARC 1P1G"
/note="Vector: PCMV SPORT6; Site 1: NotI; Site 2: SalI; library made from pooled tissue from day 11, 13, 15, 20, and 30 embryos."

ALIGNMENT SCORES:
Pred. No.: 0.00737 Length: 563
Score: 13.00 Matches: 13
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 81.25% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-3 (1-16) X BG833894 (1-563)

QY 4 GYIIeATgLeuPProleuARgSerGIyLeuGIyAla 16
|||||
DB 504 GGCATCCGGCTGCCCTGCGAAGCGGCTTGGGGGGCA 542

RESULT 5
LOCUS BP452814 616 bp mRNA linear EST 30-DEC-2003
DEFINITION BP452814 full-length enriched swine cDNA library, adult liver Sus scrofa CDNA clone LVKM10122F10 5', mRNA sequence.
ACCESSION BP452814
VERSION BP452814.1 GI:40442881
KEYWORDS EST.
SOURCE Sus scrofa (pig)
ORGANISM Sus scrofa

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
1 (bases 1 to 616)
Uenishi, H., Bguchi, T., Suzuki, K., Sawazaki, T., Toki, D., Shinkai, H., Okumura, N., Hamada, N. and Anata, T.
PEDF (pig EST Data Explorer): construction of a database for ESTs derived from porcine full-length cDNA libraries
Nucleic Acids Res. 32 (1), D484-D488 (2004)
JOURNAL CONTACT: Hirohiko Uenishi
Animal Genome Laboratory, Genome Research Department
National Institute of Agrobiological Sciences
2 Ikenodai, Tsukuba, Ibaraki 305-8602, Japan
Tel: +81-29-838-8627
Fax: +81-29-838-8627
Email: huenishi@affrc.go.jp
EST project with full-length enriched cDNA libraries carried out in Animal Genome Research Program (Japan) by National Institute of Agrobiological Sciences and STAFF-Institute
Single pass sequencing of clones derived from oligo-capped cDNA library
Vector sequences were eliminated by RepeatMasker version 2002/07/13 and crossmatch version 0.990319
Low quality bases were trimmed based on the quality values.
Location/Qualifiers
1..616
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/clone="LVKM10122F10"
/cruise_type="liver"
/dev_stage="adult"
/clone_11b="full-length enriched swine cDNA library, adult liver"

ORIGIN
Alignment Scores:
Pred. No.: 0.00798 Length: 616
Score: 13.00 Matches: 13
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 81.25% Indels: 0
DB: 5 Gaps: 0

US-10-726-967A-3 (1-16) X BP452814 (1-616)

QY 4 GYIIeATgLeuPProleuARgSerGIyLeuGIyAla 16
|||||
DB 548 GGCATCCGGCTGCCCTGCGAAGCGGCTTGGGGGGCA 566

RESULT 6
LOCUS CE200035/c 171 bp DNA linear GSS 25-SEP-2003
DEFINITION tigr-gss-dog-17000372217391 Dog Library Canis familiaris genomic.
ACCESSION CE200035
VERSION CE200035.1 GI:35355688
KEYWORDS GSS.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
REFERENCE Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Eukaryota; Eutheria; Carnivora; Fissipedia; Canidae; Canis.
1 (bases 1 to 171)
Kirkness, E.F., Batina, V., Halpern, A.L., Levy, S., Remington, K., Busch, D.B., Delcher, A.L., Pop, M., Wang, W., Frazer, C.M. and Venter, J.C.
The dog genome: survey sequencing and comparative analysis
Science 301 (5641), 1898-1903 (2003)
JOURNAL MEDLINE 22875432
PUBMED 14512627
COMMENT Contact: Kirkness EF
The Institute for Genomic Research
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive, Rockville, MD 20850, USA

Tel: 301-838-0200
Fax: 301-838-0208
Email: ekirknes@igr.org
Class: shotgun.

FEATURES

Source Location/Qualifiers

1..171
/organism="Canis familiaris"
/mol_type="Genomic DNA"
/strain="Standard Poodle"
/db_xref="taxon:9615"
/clone_lib="Dog Library"
/note="Site 1: BstXI; Libraries were prepared from peripheral blood"

ORIGIN

Alignment Scores:

Pred. No.:	0.227	Length:	171
Score:	11.00	Matches:	11
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	68.75%	Indels:	0
DB:	9	Gaps:	0

US-10-726-967A-3 (1-16) x CE200035 (1-171)

QY 4 G|Y|l|a|a|g|l|e|u|p|r|o|l|e|u|a|g|s|e|r|g|y|l|e|u|g|y 14

Db 168 GGCATCGGCTGCTGCCCTTGGCAGCGGCGTGGG 136

RESULT 7

CN692524

LOCUS

DEFINITION E0324B10-5 NIA Mouse E10.5 whole embryo cDNA library (Long) Mus

ACCESSION CN692524

VERSION CN692524.1 GI:47461272

KEYWORDS

SOURCE

ORGANISM

Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

AUTHORS

Sharov, A.A., Piao, Y., Matoba, R., Dudekula, D.B., Qian, Y., Wang, Y., Carter, M.G., Hamatani, T., Alba, K., Akutsu, H., Sharova, L., Tanaka, T.S., Kimber, W.L., Yoshikawa, T., Jaradat, S.A., Pantano, S., Nagata, R., Boheler, K.R., Taub, D., Hodess, R.J., Longo, D.L., Schlessinger, D., Keller, J., Klotz, E., Kelsoe, G., Umezawa, A., Vescevi, A.L., Rossant, J., Kunath, T., Hogan, B.L., Curci, A., D'Urso, M., Kelso, J., Hide, W. and Ko, M.S.

TITLE

JOURNAL

COMMENT

Transcriptome analysis of mouse stem cells and early embryos
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igsn.gsc.nia.nih.gov
Plate: E0324 row: B column: 10
Seq primer: M13 Reverse
High quality sequence stop: 493
POLYA=NO.

FEATURES

Source

Location/Qualifiers
1..493
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="NIA:E0324B10 IMAGE:30861237"
/tissue_type="whole embryo including extraembryonic tissues at 10.5-days postcoitum"
/dev_stage="E10.5"
/lab_host="DH10B"

ORIGIN

Alignment Scores:

Pred. No.:	5.47	Length:	493
Score:	10.00	Matches:	10
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	62.50%	Indels:	0
DB:	7	Gaps:	0

US-10-726-967A-3 (1-16) x CN692524 (1-493)

QY 4 G|Y|l|a|a|g|l|e|u|p|r|o|l|e|u|a|g|s|e|r|g|y|l|e|u|g|y 13

Db 382 GGCATCGGCTGCTGCCCTTGGCAGCGGCGTGG 411

RESULT 8

CN697484

LOCUS

DEFINITION E0394H10-5 NIA Mouse E10.5 whole embryo cDNA library (Long) Mus

ACCESSION CN697484

VERSION CN697484.1 GI:47466233

KEYWORDS

SOURCE

ORGANISM

Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

AUTHORS

Sharov, A.A., Piao, Y., Matoba, R., Dudekula, D.B., Qian, Y., Wang, Y., Carter, M.G., Hamatani, T., Alba, K., Akutsu, H., Sharova, L., Tanaka, T.S., Kimber, W.L., Yoshikawa, T., Jaradat, S.A., Pantano, S., Nagata, R., Boheler, K.R., Taub, D., Hodess, R.J., Longo, D.L., Schlessinger, D., Keller, J., Klotz, E., Kelsoe, G., Umezawa, A., Vescevi, A.L., Rossant, J., Kunath, T., Hogan, B.L., Curci, A., D'Urso, M., Kelso, J., Hide, W. and Ko, M.S.

TITLE

JOURNAL

COMMENT

Transcriptome analysis of mouse stem cells and early embryos
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igsn.gsc.nia.nih.gov
Plate: E0394 row: H column: 10
Seq primer: M13 Reverse
High quality sequence stop: 533
POLYA=NO.

FEATURES
SOURCE

Location/Qualifiers

```
1. 533
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="nabst:E0394H10-5"
/db_xref="taxon:10090"
/clone="NIA:E0394H10 IMAGE:30868029"
/tissue_type="whole embryo including extraembryonic
tissues at 10.5-days postcoitum"
/dev_stage="E10.5"
/lab_host="DH10B"
/clone_lib="NIA Mouse E10.5 whole embryo cDNA library
(long)"
/notes="Vector: pCMV-SPORT6 (Invitrogen); Site 1: SalI;
Site 2: NotI; Mouse cDNA project by the Laboratory of
Genetics, National Institute on Aging (NIA), Intramural
Research Program, NIH (http://igsun.grc.nia.nih.gov/cDNA).
This is a long-transcript enriched cDNA library (Ref.
Genome Res. 11: 1553-1558 (2001). [PMD: 11544199]). Total
RNAs were extracted from a pool of 8 embryos at 10.5-days
postcoitum. Double-stranded cDNAs were synthesized with an
Oligo(dT) primer (Invitrogen):
5'-PACTAGTCTAGATCGAGCGCGCCCTTTT-3' from
2ug of total RNA, treated with T4 DNA polymerase, and
purified by ethanol-precipitation. The cDNAs were ligated
to Lene-linker Lr-Sal4, purified by phenol/chloroform, and
separated from free linkers by Centricon 100. Then, the
cDNAs were amplified by long-range high fidelity PCR using
Ex Tag polymerase (Takara) with a primer Sal4-S. The
products were purified by phenol/chloroform and Centricon
100. The cDNAs were digested with SalI and NotI enzymes
and cloned into SalI/NotI site of pCMV-SPORT6 plasmid
vector. The DH10B E. coli host was transformed with the
ligation mixture by the standard chemical method. The
average insert size is about 3.4kb. The library was
constructed by Yulan Piao."
```

ORIGIN

Alignment Scores:

```
Score: 5.87 Length: 533
Pred. No.: 10.00 Matches: 10
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 62.50% Indels: 0
DB: 7 Gaps: 0
```

US-10-726-967A-3 (1-16) x CN697484 (1-533)

```
Qy 4 GYIIEAAGLEUPROLEUARGSGYIEU 13
    |||
Db 504 GGCATCGGCTGCCCTTCGACGGCGCTG 533
```

RESULT 9
CF903755

LOCUS 547 bp mRNA linear EST 04-NOV-2003
DEFINITION A0413D10-5 NIA Mouse Osteoblast cDNA library (Long 1) Mus musculus
CDNA clone NIA:A0413D10 IMAGE:30739245 5', mRNA sequence.

ACCESSION CF903755
VERSION CF903755.1 GI:38170804
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 547)
AUTHORS Piao, Y., Ko, N.T., Lim, M.K. and Ko, M.S.H.
TITLE Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
SUBSTRUCTURE amounts of long-transcript enriched cDNA libraries from
submicrogram amounts of total RNAs by a universal PCR amplification
method

JOURNAL Genome Res. 11 (9), 1553-1558 (2001)
MEDLINE 21429098
PubMed 11544199

COMMENT

Contact: Dawood B. Dudekula
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igsun.grc.nia.nih.gov
Plate: A0413 row: D column: 10
Seq primer: M13 Reverse
High quality sequence stop: 547
POLY-A-No.

FEATURES
SOURCE

Location/Qualifiers

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/dev_stage="K05A/A1 cells"
/lab_host="DH10B"
/clone_lib="NIA Mouse Osteoblast cDNA library (Long 1)"
/notes="Vector: pCMV-SPORT6 (Invitrogen); Site 1: SalI;
Site 2: NotI; Mouse cDNA project by the Laboratory of
Genetics, National Institute on Aging (NIA), Intramural
Research Program, NIH (http://igsun.grc.nia.nih.gov/cDNA).
This is a long-transcript enriched cDNA library (Ref.
Genome Res. 11: 1553-1558 (2001). [PMD: 11544199]). Total
RNAs were obtained from Dr. Akihito Umezawa (Keio
University School of Medicine, Japan). Double-stranded
cDNAs were synthesized with an Oligo(dT) primer
(Invitrogen):
5'-PACTAGTCTAGATCGAGCGCGCCCTTTT-3' from
2.1 ug of total RNA, treated with T4 DNA polymerase, and
purified by ethanol-precipitation. The cDNAs were ligated
to Lene-linker Lr-Sal4, purified by phenol/chloroform, and
separated from free linkers by Centricon 100. Then, the
cDNAs were amplified by long-range high fidelity PCR using
Ex Tag polymerase (Takara) with a primer Sal4-S. The
products were purified by phenol/chloroform and Centricon
100. The cDNAs were digested with SalI and NotI enzymes
and cloned into SalI/NotI site of pCMV-SPORT6 plasmid
vector. The DH10B E. coli host was transformed with the
ligation mixture by the standard chemical method. The
average insert size is about 3.0 kb. The library was
constructed by Yulan Piao."
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ORIGIN

Alignment Scores:

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Score: 6 Length: 547
Pred. No.: 10.00 Matches: 10
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 62.50% Indels: 0
DB: 7 Gaps: 0
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US-10-726-967A-3 (1-16) x CF903755 (1-547)

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Db 497 GGCATCGGCTGCCCTTCGACGGCGCTG 526
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RESULT 10
CF906581

LOCUS 563 bp mRNA linear EST 04-NOV-2003
DEFINITION A0448H07-5 NIA Mouse Osteoblast cDNA library (Long 1) Mus musculus
CDNA clone NIA:A0448H07 IMAGE:30742650 5', mRNA sequence.

ACCESSION CF906581
VERSION CF906581.1 GI:38173630
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 563)
AUTHORS Piao, Y., Ko, N.T., Lim, M.K. and Ko, M.S.H.
TITLE Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
SUBSTRUCTURE amounts of long-transcript enriched cDNA libraries from
submicrogram amounts of total RNAs by a universal PCR amplification
method

AUTHORS Piao, Y., Ko, N.T., Lim, M.K. and Ko, M.S.H.
TITLE Construction of long-transcript enriched cDNA libraries from submicrogram amounts of total RNAs by a universal PCR amplification method
JOURNAL Genome Res. 11 (9), 1553-1558 (2001)
MEDLINE 21429098
PUBMED 11544199
COMMENT Contact: Dawood B. Dudekula
Laboratory of Genetics
National Institute on Aging/National Institutes of Health
333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6820, USA
Email: cdna@igeun.igr.nia.nih.gov
Plate: A0448 row: H column: 07
Seq primer: M13 Reverse
High quality sequence stop: 563
POLY(A)=No.

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FEATURES
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Location/Qualifiers
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/clone_lib="NIA Mouse Osteoblast cDNA library (long 1)"
/notes="Vector: pCMV-SPORT6 (Invitrogen); Site 1: SalI;
Site 2: NotI; Mouse cDNA project by the Laboratory of
Genetics, National Institute on Aging (NIA), Intramural
Research Program, NIH (http://19gen.grc.nia.nih.gov/cDNA)
This is a long-transcript enriched cDNA library [Ref.
Genome Res. 11: 1553-1558 (2001). [PMID: 11544191]]. Total
RNAs were obtained from Dr. Akihito Umezawa (Keio
University School of Medicine, Japan). Double-stranded
cDNAs were synthesized with an Oligo (dT) primer
[Invitrogen]:
5'-GAGCTAGTCTTACAGATCGACGCGCGCCCTTTT-3' from
5'-1g of total RNA, treated with T4 DNA polymerase, and
purified by ethanol-precipitation. The cDNAs were ligated
to lone-linker II-SalI, purified by phenol/chloroform, and
separated from free linkers by Centricon 100. Then, the
cDNAs were amplified by long-range high fidelity PCR using
Ex Taq polymerase (Takara) with a primer SalI-S. The
products were purified by phenol/chloroform and Centricon
100. The cDNAs were digested with SalI and NotI enzymes
and cloned into SalI/NotI site of pCMV-SPORT6 plasmid
vector. The DH10B E. coli host was transformed with the
ligation mixture by the standard chemical method. The
average insert size is about 3.0 Kd. The library was
constructed by Yulan Piao."

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ORIGIN					
Alignment Scores:					
Pred. No.:	6.16	Length:	563		
Score:	10.00	Matches:	10		
Percent Similarity:	100.00%	Conservative:	0		
Best local Similarity:	100.00%	Mismatches:	0		
Query Match:	62.50%	Indels:	0		
DB:	7	Gaps:	0		
US-10-726-967A-3 (1-16) x CF906581 (1-563)					
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Dd	497	GGCATCCGCGCTGCCCTTCGGAGCGGCTCG	526		
RESULT 11					
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DEFINITION	B0839F04-5 NIA Mouse Newborn Kidney cDNA Library	(Long 1)	Mus		
ACCESSION	CF171218	mMusculus cDNA clone NIA:B0839F04 IMAGE:30471231	5'	mRNA sequence.	

VERSION	CF171218.1	GI:33280767
KEYWORDS	EST.	
SOURCE	Mus musculus	(house mouse)
ORGANISM	Mus musculus	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.	
AUTHORS	1 (bases 1 to 576)	
TITLE	Piao, Y., Ko, N.T., Lim, M.K. and Ko, M.S.H.	
JOURNAL	Construction of long-transcript enriched cDNA libraries from	
MEDLINE	submicrogram amounts of total RNAs by a universal PCR amplification	
PUBMED	method	
COMMENT	Genome Res. 11 (9), 1553-1558 (2001)	
	21429098	
	11544139	
	Contact: Dawood B. Dudekula	
	Laboratory of Genetics	
	National Institute on Aging/National Institutes of Health	
	333 Cassell Drive, Suite 4000, Baltimore, MD 21224-6620, USA	
	Email: cdaa@gsun.grc.nia.nih.gov	
	Phone: 80839 row: F column: 04	
	Seq primer: M13 Reverse	
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FEATURES
Source

Location/Qualifiers
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/strain="C57BL/6J"
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1)"
/note="Vector: pCMV-SPORT6 (Invitrogen); Site_1: SalI;
Site_2: NotI; Mouse cDNA project by the Laboratory of
Genetics, National Institute on Aging (NIA), Intramural
Research Program, NIH (http://lgsun.grc.nia.nih.gov/cDNA).
In brief, double-stranded cDNAs were synthesized with an
Oligo(dt) primer (Invitrogen:
5'-GACATGCTTCAGATCGCGAGCGCCGCTTTT-TTTT-3') from
26 ug of total RNA, treated with T4 DNA polymerase, and
purified by ethanol-precipitation. The cDNAs were ligated
to lone-linker Lp-Sal-I, purified by phenol/chloroform, and
separated from free linkers by Centricon 100. Then, the
cDNAs were amplified by long-range high fidelity PCR using
Ex Taq polymerase (Takara) with a primer Sal-I-S. The
products were purified by phenol/chloroform and Centricon
100. The cDNAs were digested with SalI and NotI enzymes
and cloned into SalI/NotI site of pCMV-Sports plasmid
vector. The DH10B E. coli host was transformed with the
ligation mixture by the standard chemical method. The
average insert size is about 3.0 kb. The library was
constructed by Yulan Piao."

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	ALIGNMENT SCORES:	
Pred. No.:	6, 29	576
Score:	10, .00	10
Percent Similarity:	100.00%	
Best Local Similarity:	Conservative:	0
Query Match:	Mismatches:	0
	Indels:	0
DB:	Gaps:	0
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Dd	497 GGCACTGGCGTCCGCCCTTCAGACGCACCNG 526	
RESULT 12		
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DEFINITION BY173879 RIKEN full-length enriched, 16 days embryo head Mus
musculus cDNA clone 4122401C04 5', mRNA sequence.
ACCESSION BY173879
VERSION BY173879.1 GI:27126044
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus (house mouse)
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 600)
Oikazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Otsu, N., Saito, R., Suzuki, H., Yamanaka, I.,
Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A.,
Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D.P., Bult, C.,
Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuda, H.,
Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Brusic, V.,
Chochia, C., Corbani, L.E., Cousins, S., Dalla, E., Dragan, T.A.,
Fletcher, C.F., Forrest, A., Fraser, K.S., Gaasterland, T.,
Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S.,
Guelinckich, S., Hirokawa, N., Jackson, I.J., Jarvis, E.D., Kanai, A.,
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Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M.,
Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y.,
Wells, C., Wilming, L.G., Wymshew-Boris, A., Yanagisawa, M., Yang, I.,
Yang, L., Yuan, Z., Zavalan, M., Zhu, Y., Zimmer, A., Carninci, P.,
Hayatsu, N., Hirozane-Kishikawa, T., Kono, H., Nakamura, M.,
Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y.,
Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K.,
Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E.S.,
Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
22354683
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@sc.riken.jp, URL: http://genome.gsc.riken.jp/
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P.,
Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F.,
Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y.,
Kondo, S., Kono, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M.,
Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N.,
Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M.,
Takeda, Y., Waki, K., Watanishi, A., Muramatsu, M. and Hayashizaki, Y.
Direct Submision
Computational Analysis of Full-length Mouse cDNAs Compared with
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system-384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.

FEATURES
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head"
/note="Site 1: SalI; Site 2: BamHI. cDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in
RIKEN. Division of Experimental Animal Research in Riken
contributed to prepare mouse tissues. 1st strand cDNA was
primed with a primer [5'
GAGAGAGAGAGATCCAGAGCTCTTTTCTTTTCTTTTNN 3'], cDNA was
transcribed by using trehalose thermo-activated reverse
transcriptase and subsequently enriched for full-length by
cap-trapper. Second strand cDNA was prepared with the
primer adapter of sequence [5'
GAGAGAGATCTCGAGCTTATTAATTAATTCCTCCCCCCCC 3']. cDNA
was cloned into the xhoI and BamHI sites. Vector: a
modified pBluescript KS(+) after bulk excision from lambda
FLC I"

ORIGIN
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Pred. No.: 6.52 Length: 600
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Query Match: 62.50% Indels: 0
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QY 4 Gly11eartgleuProleuArgSerGlylen 13
DB 498 GGCATCCGCGCTGCCCCCTTCGACGCGCTG 527
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ACCESSION BB644736
VERSION BB644736.1 GI:16479273
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus (house mouse)
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 619)
Arakawa, T., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T.,
Hara, A., Hiramoto, K., Hori, F., Ishii, Y., Ito, M., Kawai, J.,
Kono, H., Konda, M., Koya, S., Matsuyama, T., Miyazaki, A., Nomura, K.,
Ohno, M., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F.,
Takeda, Y., Tanaka, T., Toya, T., Muramatsu, M. and Hayashizaki, Y.
RIKEN Mouse ESTs (Arakawa, T., et al. 2001)
Unpublished (2001)
TITLE
JOURNAL
COMMENT
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic

LOCUS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE	ORGANISM	REFERENCE	AUTHORS
B6392244	639 bp mRNA linear EST 26-OCT-2001	B6392244	RIKEN full-length enriched, adult male hypothalamus Mus musculus cDNA clone A230019C07 5', mRNA sequence.	B6392244	B6392244.1 GI:16468863	Mus musculus (house mouse)	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.	EST.
TITLE	JOURNAL	COMMENT						
RIKEN Mouse ESTs (Arakawa,T., et al. 2001) Unpublished (2001)	Contact: Yoshihide Hayashizaki Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute The Institute of Physical and Chemical Research (RIKEN) 1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan Tel.: 81-45-503-9222 Fax: 81-45-503-9216 Email: genome-res@gsr.riken.jp, URL:http://genome.gsc.riken.jp/							
Normalisation and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)	wagi,K., Fujiwara,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E., Watabiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T., Matsura,S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kita,A., and Hayashizaki,Y.							
RIKEN integrated sequence analysis (RISA) system-384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)	Kono,H., Pukushihashi,Y., Shibata,K., Itoh,M., Carninci,P., Sugahara,Y. and Hayashizaki,Y.							
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)	Kondo,S., Shingawa,A., Saito,T., Kiyosawa,H., Yamanae,I., Aizawa,K., Fukuda,S., Hara,A., Itoh,M., Kawai,J., Shibata,K. and Hayashizaki,Y.							
Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences. Mamm. Genome. 12, 673-677 (2001)	Please visit our web site (http://genome.gsc.riken.go.jp) for further details.							
e mouse tissues.	Location/Qualifiers							
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ORIGIN
Alignment Scores:
Pred. No.:          6 89
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Best Local Similarity: 100.00%
Query Match:        62.50%
DB:                  2
US-10-726-967A-3 (1-16) x BB632244 (1-639)
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Db      518 GGCATCGGCGCTGCCCTCCGACACGGCCCTG 547
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Job time : 3150 secs

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GAGAGAAGAAAGATCCCAAGACTCTTTTTTTTTTTTTNN 3'], cDNA was prepared by using trihalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 20.0 and subtraction to Rot = 459.0. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAAGATTCCTCAGTAATAATTATGCCCCCCCCCCC 3']. cDNA was cleaved with XhoI and BamHI. Vector: a modified p Bluescript KS(+) after bulk excision from Lambda

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - nucleic search, using frame_plus.p2n model

Run on: July 27, 2005, 16:41:37 ; Search time 1935 Seconds
(without alignments)
701.161 Million cell updates/sec

Title: US-10-726-967A-52

Perfect score: 144
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Scoring table:
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Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4708233 seqs, 24227607955 residues
Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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2	144	100.0	1278	6	BD235898 Alzheimer
3	144	100.0	1278	6	AR224104 Sequence
4	144	100.0	1278	6	AR269235 Sequence

5	144	100.0	1278	6	AR478790	AR478790	Sequence
6	144	100.0	1278	6	AR487356	AR487356	Sequence
7	144	100.0	1278	6	AR531996	AR531996	Sequence
8	144	100.0	1278	6	AR540897	AR540897	Sequence
9	144	100.0	1278	6	AR560107	AR560107	Sequence
10	144	100.0	1278	6	AR105409	AR105409	Sequence
11	144	100.0	1278	6	AR224122	AR224122	Sequence
12	144	100.0	1287	6	AR269253	AR269253	Sequence
13	144	100.0	1287	6	AR478808	AR478808	Sequence
14	144	100.0	1287	6	AR487374	AR487374	Sequence
15	144	100.0	1287	6	AR532014	AR532014	Sequence
16	144	100.0	1287	6	AR540915	AR540915	Sequence
17	144	100.0	1287	6	AR560125	AR560125	Sequence
18	144	100.0	1287	6	AX105432	AX105432	Sequence
19	144	100.0	1287	6	AX573870	AX573870	Sequence
20	144	100.0	1287	6	BD235897	BD235897	Sequence
21	144	100.0	1302	6	AR224103	AR224103	Sequence
22	144	100.0	1302	6	AR269234	AR269234	Sequence
23	144	100.0	1302	6	AR478789	AR478789	Sequence
24	144	100.0	1302	6	AR487355	AR487355	Sequence
25	144	100.0	1302	6	AR531995	AR531995	Sequence
26	144	100.0	1302	6	AR540896	AR540896	Sequence
27	144	100.0	1302	6	AR560106	AR560106	Sequence
28	144	100.0	1302	6	AX105407	AX105407	Sequence
29	144	100.0	1302	6	AX573845	AX573845	Sequence
30	144	100.0	1302	6	AX700454	AX700454	Sequence
31	144	100.0	1305	6	AR224123	AR224123	Sequence
32	144	100.0	1305	6	AR269254	AR269254	Sequence
33	144	100.0	1305	6	AR478809	AR478809	Sequence
34	144	100.0	1305	6	AR487375	AR487375	Sequence
35	144	100.0	1305	6	AR532015	AR532015	Sequence
36	144	100.0	1305	6	AR540916	AR540916	Sequence
37	144	100.0	1305	6	AR560126	AR560126	Sequence
38	144	100.0	1305	6	AX105434	AX105434	Sequence
39	144	100.0	1305	6	AX573872	AX573872	Sequence
40	144	100.0	1305	6	AB050438	AB050438	Homo sapi
41	144	100.0	1333	9	BD235895	BD235895	Alzheimer
42	144	100.0	1341	6	AR224101	AR224101	Sequence
43	144	100.0	1341	6	AR269232	AR269232	Sequence
44	144	100.0	1341	6	AR478787	AR478787	Sequence
45	144	100.0	1341	6			

ALIGNMENTS

RESULT 1	AB089958	517 bp	mRNA	linear	PRI 19-AUG-2003
LOCUS	AB089958				
DEFINITION	Homo sapiens BACE mRNA for beta-site APP cleaving enzyme isoform I-127, complete cds.				
ACCESSION	AB089958				
VERSION	AB089958.1	GI:34014375			
KEYWORDS					
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	1				
AUTHORS	Tanahashi, H.				
TITLE	A novel alternatively spliced isoform of BACE, I-127 induced by cycloheximide treatment				
JOURNAL	Unpublished				
REFERENCE	2	(bases 1 to 517)			
AUTHORS	Tanahashi, H.				
TITLE	Direct Subregion				
JOURNAL	Submitted (17-AUG-2002) Hiroshi Tanahashi, National Institute of Neuroscience, Division of Demyelinating Disease and Aging; 4-1-1 Ogawahigashi, Kodaira, Tokyo 187-8502, Japan				
FEATURES	(E-mail:tanahashicnp.go.jp, Tel:81-042-341-2711 (ex. 5163), Fax:81-042-346-1747)				
source	Location/Qualifiers				
	1..517				
	/organism="Homo sapiens"				

/mol_type="mRNA"
/db_xref="taxon:9606"
/chromosome="11"
/map="11q23.2-23.3"
/cell_line="human neuroblastoma SH-SY5Y"
/note="I-127 is induced by cycloheximide treatment.
Alternative splicing of the RNA occurs at an internal
donor in exon 3."
1..517
/gene="BACE"
1..384
/gene="BACE"
/codon_start=1
/product="beta-site APP cleaving enzyme isoform I-127"
/protein_id="BAC81826.1"
/db_xref="GI:34014376"
/translation="MAQALPMLLMGAGVLPNHTQHGIRLPPLRSGLGAPLGRLP
RRTDEPFEPRGRGSFEVNDLGRKSGGGYVEMTVGSPPTLNILVDTGSSNFAVG
AAPHPFLHRYQRQLSSTYRDLRKA"

ORIGIN

Alignment Scores:
Pred. No.: 1,13e-14 Length: 517
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967A-52 (1-28) x AB089958 (1-517)

QY 1 G|Y|T|Y|T|V|A|G|U|E|T|H|T|V|A|G|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|N|I|L|E|U|V|A|L|A|S|P|20
Db 220 G|G|C|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|G|C|A|G|A|C|C|C|T|C|A|A|C|T|C|T|G|T|G|A|T|279

QY 21 T|H|G|Y|S|E|R|S|E|R|A|P|H|E|A|L|A|V|A|L|28
Db 280 A|C|A|G|G|C|A|G|C|A|G|T|T|T|G|C|A|G|T|G|303

RESULT 2
BD235898
LOCUS BD235898 1278 bp DNA linear PAT 17-JUL-2003
DEFINITION Alzheimer's disease secretase.
ACCESSION BD235898.1 GI:33045668
VERSION BD235898.1
KEYWORDS JP 2002526081-A/14.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 1278)
Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
Yan,R.
Alzheimer's disease secretase
Patent: JP 2002526081-A 14 20-AUG-2002;
PHARMACIA AND UPJOHN CO
OS Homo sapiens (human)
PN JP 2002526081-A/14
PD 20-AUG-2002
PR 23-SEP-1999 JP 2000574268
PF 24-SEP-1998 US 60/101594
PI MARK E GURNEY, MICHAEL JEROME BIENKOWSKI, ROBERT LEROY PI
HEINRICHSON,
PI LUIS A PARODI, RIOTANG VAN
PC C12N15/09,A61K45/00,A61P25/28,C07K14/47,C07K16/18,C12N1/15, PC
C12N1/19,
PC C12N1/21,C12N5/10,C12N9/64,C12P21/02,C12P21/08,C12Q1/37,G01N33/ PC
15,
G01N33/50//C12N1/21,C12P1/19,C12N5/00,C12N5/00 CC
Alzheimer's disease secretase
FH Key Location/Qualifiers
FT source 1..1278

FEATURES FT
source Location/Qualifiers
1..1278
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"

ORIGIN

Alignment Scores:
Pred. No.: 3.18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR224104 (1-1278)

QY 1 G|Y|T|Y|T|V|A|G|U|E|T|H|T|V|A|G|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|N|I|L|E|U|V|A|L|A|S|P|20
Db 136 G|G|C|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|G|C|A|G|A|C|C|C|T|C|A|A|C|T|C|T|G|T|G|A|T|195

QY 21 T|H|G|Y|S|E|R|S|E|R|A|P|H|E|A|L|A|V|A|L|28
Db 196 A|C|A|G|G|C|A|G|C|A|G|T|T|T|G|C|A|G|T|G|219

RESULT 3
AR224104
LOCUS AR224104 1278 bp DNA linear PAT 26-SEP-2002
DEFINITION Sequence 27 from patent US 6440698.
ACCESSION AR224104
VERSION AR224104.1 GI:23332764
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
Yan,R.
Alzheimer's disease secretase, APP substrates therefor, and uses
therefor
Patent: US 6440698-A 27 27-AUG-2002;
LOCATION/Qualifiers
1..1278
source
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Alignment Scores:
Pred. No.: 3.18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR224104 (1-1278)

QY 1 G|Y|T|Y|T|V|A|G|U|E|T|H|T|V|A|G|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|N|I|L|E|U|V|A|L|A|S|P|20
Db 136 G|G|C|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|G|C|A|G|A|C|C|C|T|C|A|A|C|T|C|T|G|T|G|A|T|195

QY 21 T|H|G|Y|S|E|R|S|E|R|A|P|H|E|A|L|A|V|A|L|28
Db 196 A|C|A|G|G|C|A|G|C|A|G|T|T|T|G|C|A|G|T|G|219

RESULT 4
AR269235
LOCUS AR269235 1278 bp DNA linear PAT 10-APR-2003
DEFINITION Sequence 27 from patent US 6500667.
ACCESSION AR269235
VERSION AR269235.1 GI:29700203
KEYWORDS

SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrikson,R.L., Parodi,L.A. and Yan,R.
TITLE Aspartyl protease 2 (Asp2) antisense oligonucleotides
JOURNAL Patent: US 6500667-A 27 31-DEC-2002;
FEATURES Location/Qualifiers
source 1..1278
/organism="unknown"
/mol_type="genomic DNA"

Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR269235 (1-1278)

QY 1 G|Y|Y|Y|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|N|I|L|E|U|V|A|A|P 20
DB 136 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|G|C|C|C|C|C|C|C|A|G|A|C|G|C|T|C|A|A|C|A|T|C|C|T|G|G|G|A|T 195

QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|A|V|A| 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|G|C|A|G|T|G 219

RESULT 5
AR478790
LOCUS AR478790 1278 bp DNA linear PAT 14-MAY-2004
DEFINITION Sequence 27 from patent US 669671.
ACCESSION AR478790
VERSION AR478790.1 GI:47237510
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrikson,R.L., Parodi,L.A. and Yan,R.
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses therefor
JOURNAL Patent: US 669671-A 27 02-MAR-2004;
FEATURES Location/Qualifiers
source 1..1278
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR478790 (1-1278)

QY 1 G|Y|Y|Y|Y|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|N|I|L|E|U|V|A|A|P 20
DB 136 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|G|C|C|C|C|C|C|C|A|G|A|C|G|C|T|C|A|A|C|A|T|C|C|T|G|G|G|A|T 195

QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|A|V|A| 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|G|C|A|G|T|G 219

RESULT 6

AR487356
LOCUS AR487356 1278 bp DNA linear PAT 14-MAY-2004
DEFINITION Sequence 27 from patent US 6706485.
ACCESSION AR487356
VERSION AR487356.1 GI:47252454
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrikson,R.L., Parodi,L.A. and Yan,R.
TITLE Method of identifying agents that inhibit APP processing activity
JOURNAL Patent: US 6706485-A 27 16-MAR-2004;
FEATURES Location/Qualifiers
source 1..1278
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR487356 (1-1278)

QY 1 G|Y|Y|Y|Y|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|N|I|L|E|U|V|A|A|P 20
DB 136 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|G|C|C|C|C|C|C|C|A|G|A|C|G|C|T|C|A|A|C|A|T|C|C|T|G|G|G|A|T 195

QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|A|V|A| 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|G|C|A|G|T|G 219

RESULT 7
AR531996
LOCUS AR531996 1278 bp DNA linear PAT 08-OCT-2004
DEFINITION Sequence 27 from patent US 6727074.
ACCESSION AR531996
VERSION AR531996.1 GI:53920530
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrikson,R.L., Parodi,L.A. and Yan,R.
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses therefor
JOURNAL Patent: US 6727074-A 27 27-APR-2004;
FEATURES Location/Qualifiers
source 1..1278
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR531996 (1-1278)

QY 1 G|Y|Y|Y|Y|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|N|I|L|E|U|V|A|A|P 20
DB 136 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|G|C|C|C|C|C|C|C|A|G|A|C|G|C|T|C|A|A|C|A|T|C|C|T|G|G|G|A|T 195

QY 21 ThrGlySerSerAanPheAlaVal 28
Db 196 ACAGGACAGCACTTTCAGCTG 219

RESULT 8
ARS40897
LOCUS ARS40897 1278 bp DNA linear PAT 08-OCT-2004
DEFINITION Sequence 27 from patent US 6737510.
ACCESSION ARS40897
VERSION ARS40897.1 GI:53932410
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and Yan,R.
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses thereof
JOURNAL Patent: US 6737510-A 27 18-MAY-2004;
FEATURES
source
Location/Qualifiers
1..1278
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: Gaps: 0

US-10-726-967A-52 (1-28) x ARS40897 (1-1278)

QY 1 GlyTyrTyrValGluMetThrValGlySerProProGlnThrLeuAanlleuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGACCCCCCGACAGCCTCAACATCTGTGGAT 195

QY 21 ThrGlySerSerAanPheAlaVal 28
Db 196 ACAGGACAGCACTTTCAGCTG 219

RESULT 9
ARS60107
LOCUS ARS60107 1278 bp DNA linear PAT 08-OCT-2004
DEFINITION Sequence 27 from patent US 6753163.
ACCESSION ARS60107
VERSION ARS60107.1 GI:53970474
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 1278)
AUTHORS Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and Yan,R.
TITLE Alzheimer's disease secretase, APP substrates therefor, and uses thereof
JOURNAL Patent: US 6753163-A 27 22-JUN-2004;
FEATURES
source
Location/Qualifiers
1..1278
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN
Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: Gaps: 0

DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x ARS60107 (1-1278)

QY 1 GlyTyrTyrValGluMetThrValGlySerProProGlnThrLeuAanlleuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGACCCCCCGACAGCCTCAACATCTGTGGAT 195

QY 21 ThrGlySerSerAanPheAlaVal 28
Db 196 ACAGGACAGCACTTTCAGCTG 219

RESULT 10
AX105409
LOCUS AX105409 1278 bp DNA linear PAT 30-APR-2001
DEFINITION Sequence 27 from Patent WO0123533.
ACCESSION AX105409
VERSION AX105409.1 GI:13921523
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
REFERENCE 1
AUTHORS Gurney,M. and Bienkowski,M.J.
TITLE Alzheimer's disease secretase, app substrates therefor, and uses thereof
JOURNAL Patent: WO 0123533-A 27 05-APR-2001;
FEATURES
source
Location/Qualifiers
1..1278
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
Alignment Scores:
Pred. No.: 3,18e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: Gaps: 0

US-10-726-967A-52 (1-28) x AX105409 (1-1278)

QY 1 GlyTyrTyrValGluMetThrValGlySerProProGlnThrLeuAanlleuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGACCCCCCGACAGCCTCAACATCTGTGGAT 195

QY 21 ThrGlySerSerAanPheAlaVal 28
Db 196 ACAGGACAGCACTTTCAGCTG 219

RESULT 11
AX573847
LOCUS AX573847 1278 bp DNA linear PAT 07-JAN-2003
DEFINITION Sequence 27 from Patent EP1249498.
ACCESSION AX573847
VERSION AX573847.1 GI:27551489
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
REFERENCE 1
AUTHORS Gurney,M. and Bienkowski,M.J.
TITLE Alzheimer's disease secretase, app substrates therefor, and uses thereof
JOURNAL Patent: EP 1249498-A 27 16-OCT-2002;
FEATURES
source
Location/Qualifiers
1..1278

ORGANISM	Unknown. Unclassified.
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AR487374	1287 bp	DNA	linear	PAT 14-MAY-2004
LOCUS				
AB487374				

AR487374	1287 bp	DNA	linear	PAT 14-MAY-2004
LOCUS				
AB487374				

ACCESSION AR487374
 VERSION AR487374.1 GI:47252472
 KEYWORDS
 SOURCE
 ORGANISM Unknown.
 Unclassified.
 REFERENCE 1 (bases 1 to 1287)
 Gurney,M.E., Bienkowski,M.J., Heinrichson,R.L., Parodi,L.A. and
 Yan,R.
 TITLE Method of identifying agents that inhibit APP processing activity
 JOURNAL Patent: US 6706485-A 50 16-MAR-2004;
 FEATURES Location/Qualifiers
 source 1..1287
 /organism="Unknown"
 /mol_type="genomic DNA"

ORIGIN
 Alignment Scores:
 Pred. No.: 3.2e-14 Length: 1287
 Score: 144.00 Matches: 28
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x AR487374 (1-1287)

QY 1 G|Y|Y|Y|Y|V|A|G|U|Me|Th|r|V|A|G|Y|S|e|P|P|O|G|I|N|Th|L|e|U|A|n|L|e|U|V|A|A|p 20
 |||||
 Db 220 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|G|C|A|G|A|C|C|T|C|A|C|A|T|C|T|G|G|T|G|A|T 279
 QY 21 T|h|r|G|Y|S|e|S|e|r|A|s|n|P|h|e|A|A|V|A| 28
 |||||
 Db 280 A|C|A|G|G|C|A|G|C|A|T|T|G|C|A|G|T|G 303

Search completed: July 27, 2005, 18:50:18
 Job time : 1939 secs

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OM protein - nucleic search, using frame_plus.p2n model

Run on: July 27, 2005, 12:14:54 ; Search time 437 Seconds
(without alignments)
379.297 Million cell updates/sec

Title: US-10-726-967A-52

Sequence: 1 GYVEMTVGSPPTINILVDITGSSNFAV 28

Scoring table:

BLOSUM62			
Xgapop 10.0	Xgapext 0.5		
Ygapop 10.0	Ygapext 0.5		
Fgapop 6.0	Fgapext 7.0		
Delop 6.0	Delext 7.0		

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Command line parameters:

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-Q/cgn2.1/USPTO.spool.p/US10726567/funat.26072005.130733.6129/app.query.fasta.1.199
-DB=N Geneseg.16Dec04 -QPMF=fastap -SUFFIX=p2n.rng -MINMATCH=0.1 -LOOPEXT=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=biosum62 -TRANS=humand0.cdi
-LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15
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-NO MAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSBLOCK=100 -LONGLOG
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FAPEXT=7 -XGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

N_Geneseg.16Dec04:*
1: geneseg1980s:*
2: geneseg1990s:*
3: geneseg2000s:*
4: geneseg2001s:*
5: geneseg2001bs:*
6: geneseg2002as:*
7: geneseg2002bs:*
8: geneseg2003as:*
9: geneseg2003bs:*
10: geneseg2003cs:*
11: geneseg2003ds:*
12: geneseg2004as:*
13: geneseg2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	144	100.0	1278	3	AAAI5677 Human Asp
2	144	100.0	1278	4	AAAI1714 DNA encod
3	144	100.0	1278	4	AAAI1787 T7-Caspas
4	144	100.0	1278	4	AAAI3033 T7-Caspas
5	144	100.0	1278	4	AAAI06751 T7-Caspas

6	144	100.0	1278	4	AAAI529	AAAI529 T7-Caspas
7	144	100.0	1278	6	ABLS2469	ABLS2469 T7-Caspas
8	144	100.0	1278	12	ADJ94339	ADJ94339 Human T7-
9	144	100.0	1278	12	ADOS0435	ADOS0435 T7-Caspas
10	144	100.0	1278	13	ADR75348	ADR75348 T7-Caspas
11	144	100.0	1287	4	AAAI7895	AAAI7895 Human-Asp
12	144	100.0	1287	4	AAAI13276	AAAI13276 Human-Asp
13	144	100.0	1287	4	AAAI06768	AAAI06768 Human-Asp
14	144	100.0	1287	4	AAAI1547	AAAI1547 Human CDN
15	144	100.0	1287	6	ABLS2487	ABLS2487 Human Asp
16	144	100.0	1287	12	ADJ94362	ADJ94362 Human-pro
17	144	100.0	1287	12	ADOS0458	ADOS0458 Human Asp
18	144	100.0	1287	13	ADR75371	ADR75371 Human Asp
19	144	100.0	1302	3	AAAI5670	AAAI5670 Human-pro
20	144	100.0	1302	4	AAAI1773	AAAI1773 DNA encod
21	144	100.0	1302	4	AAAI7876	AAAI7876 Human-pro
22	144	100.0	1302	4	AAAI3032	AAAI3032 Human-pro
23	144	100.0	1302	4	AAAI06750	AAAI06750 Human-pro
24	144	100.0	1302	4	AAAI1528	AAAI1528 Human CDN
25	144	100.0	1302	6	ABLS2468	ABLS2468 Human-pro
26	144	100.0	1302	12	ADJ94337	ADJ94337 Human-pro
27	144	100.0	1302	12	ADOS0433	ADOS0433 Human-pro
28	144	100.0	1302	13	ADR75346	ADR75346 Human-pro
29	144	100.0	1305	4	AAAI1733	AAAI1733 DNA encod
30	144	100.0	1305	4	AAAI7896	AAAI7896 Human-Asp
31	144	100.0	1305	4	AAAI3277	AAAI3277 Human-Asp
32	144	100.0	1305	4	AAAI06769	AAAI06769 Human-Asp
33	144	100.0	1305	4	AAAI1548	AAAI1548 Human CDN
34	144	100.0	1305	6	ABLS2468	ABLS2468 Human Asp
35	144	100.0	1305	12	ADJ94364	ADJ94364 Human-pro
36	144	100.0	1305	12	ADOS0460	ADOS0460 Human Asp
37	144	100.0	1305	13	ADR75373	ADR75373 Human Asp
38	144	100.0	1341	3	AAAI5668	AAAI5668 T7-Caspas
39	144	100.0	1341	4	AAAI1771	AAAI1771 DNA encod
40	144	100.0	1341	4	AAAI7874	AAAI7874 T7-Human-
41	144	100.0	1341	4	AAAI3030	AAAI3030 T7-Human-
42	144	100.0	1341	4	AAAI06748	AAAI06748 T7-Human-
43	144	100.0	1341	4	AAAI1526	AAAI1526 Human CDN
44	144	100.0	1341	6	ABLS2466	ABLS2466 T7-human-
45	144	100.0	1341	12	ADJ94333	ADJ94333 Human CDN

ALIGNMENTS

RESULT 1
ID AAAI5677 standard; DNA; 1278 BP.
XX
AC AAAI5677;
XX
DT 03-AUG-2000 (first entry)
XX
DE Human Asp2 nucleotide sequence containing proteolytic cleavage site.
XX
KW Aspartyl protease; aspartase; amyloid precursor protein; APP; Asp 2;
KW Alzheimer's disease; beta secretase site; ss.
XX
OS Homo sapiens.
XX
PN WO200017369-A2.
XX
PD 30-MAR-2000.
XX
PF 23-SEP-1999; 99WO-US020881.
XX
PR 24-SEP-1998; 98US-0101594P.
XX
PA (PHAA) PHARMACIA & UPJOHN CO.
XX
PI Gurney ME, Bienkowski MJ, Heintzson RL, Parodi LA, Yan R;
XX WPI; 2000-303209/26.
XX P-PSDB; AAY88437.

XX New enzyme designated human aspartase useful in research into Alzheimer's
PT Disease is capable of cleaving amyloid protein precursor at the beta
PT secretase site to produce amyloid beta peptide.
XX

Example 9; Page 165; 183pp; English.

XX This sequence represents a modified version of the human aspartase 2
CC (Asp2) nucleotide sequence. The sequence is used in the bacterial
CC expression of human Asp2L. The invention relates to a protease (e.g.
CC Asp2) capable of cleaving the beta secretase site of amyloid precursor
CC protein (APP). The protease contains a sequence encoding the amino acid
CC sequence DTG and a sequence encoding DSG or DTG separated by 100-300
CC amino acids. When mutated the APP gene causes an autosomal dominant form
CC of Alzheimer's disease. APP localises to the cell surface membrane and
CC have a single C-terminal transmembrane domain. Proteolytic processing of
CC APP produces the amyloid beta protein, which is possibly very important
CC in Alzheimer's disease. The invention includes a nucleotide sequence
CC encoding the protease, a vector containing the nucleotide sequence, and a
CC cell line comprising the vector. Methods for screening for inhibitors of
CC beta secretase activity are also given in the invention. The human
CC aspartase protein and nucleotide sequences and the methods for
CC identifying inhibitors of the protease, are useful in the treatment of
CC and research in to Alzheimer's disease
XX

SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 4.8e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-10-726-967A-52 (1-28) x AAS11714 (1-1278)

QY 1 G|Y|T|T|T|V|A|G|L|W|E|T|H|V|A|G|L|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|N|I|L|E|U|V|A|A|S|P 20
DB 136 G|G|C|T|A|C|T|A|G|T|G|A|G|T|G|A|C|C|G|G|G|C|A|C|C|C|C|C|G|C|A|G|C|G|C|T|C|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|H|G|I|S|E|R|S|E|R|A|S|P|H|E|A|I|A|V|A| 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|T|C|A|G|T|G 219

RESULT 2

AAS11714
ID AAS11714 standard; DNA; 1278 BP.
XX

AC AAS11714;
XX

DT 11-SEP-2003 (revised)
XX
DT 24-OCT-2001 (first entry)
XX

DE DNA encoding T7-caspase-caespase 8-human aspartyl protease 2a deltaTM.

XX Human; aspartyl protease 1; Asp-1; neutrotropic; neutrotrophic;
KW aspartyl protease 2; Asp2; amyloid protein precursor; APP;
KW beta-secretase; Alzheimer's disease; ds.
XX

OS Homo sapiens.
OS Enterobacteria phage T7.
XX

XX Key Location/Qualifiers
FH CDS 1..1278
FT /*tag= a
FT /product= "T7-caspase-caespase 8-Aspartyl protease-2a
FT delta TM"
XX

MO200149097-A2.
XX

12-JUL-2001.
XX

PF 09-MAY-2001; 2001WO-IB000797.
XX
XX 09-MAY-2001; 2001WO-IB000797.
XX

PA (BIEN/) BIENKOWSKI M J.
PA (GURNEY) GURNEY M E.
PA (HEIN/) HEINRIKSON R L.
PA (PARO/) PARODI L A.
PA (YANR/) YAN R.
XX

PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
XX

DR WPI, 2001-502548/55.
XX
DR P-PSDB; AAN07214.
XX

PT Novel purified polypeptide comprising fragment of mammalian aspartyl
PT protease 2, lacking Asp2 transmembrane domain and retaining beta
PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
XX activity.
XX

Example 9; Page 158; 185pp; English.

XX The invention relates to a novel purified polypeptide comprising a
CC fragment of mammalian aspartyl protease 2 (Asp2) protein which lacks the
CC Asp2 transmembrane domain and the Asp2 protein, and where the polypeptide
CC and the fragment retain the beta-secretase activity of the mammalian Asp2
CC protein. Also included is an isoform of amyloid protein precursor (APP)
CC comprising the amino acid sequence of a APP or its fragment containing an
CC APP cleavage site recognizable by a mammalian beta-secretase, and further
CC comprising two lysine residues at the carboxyl terminus of the amino acid
CC sequence of the mammalian APP or APP fragment. The polypeptides are used
CC for assaying for modulators of beta-secretase activity. Identifying
CC agents that inhibit the APP processing activity of human Asp2 aspartyl
CC protease (Hu-Asp2); identifying agents that modulate the activity of Asp2
CC; and for reducing cellular production of amyloid beta (Abeta) from APP.
CC Agents identified by the above methods are useful for treating
CC Alzheimer's disease; and for identifying modulators of amyloid-beta
CC (Abeta) peptide production, for use in designing therapeutics for the
CC treatment or prevention of Alzheimer's disease. Probes and primers
CC derived from Asp nucleic acid sequences are useful for detecting Hu-Asp
CC nucleic acids in vitro assays and in Northern and Southern blots. The
CC present sequence represents the coding sequence of T7-caspase-caespase 8-
CC human-Asp-2a delta TM construct which has a T7 tag, a caspase 8 leader
CC sequence and cleavage site, and lacks the transmembrane domain. This
CC construct was used for bacterial expression and purification of human
CC Asp2a. (Updated on 11-SEP-2003 to standardise OS field)
XX

SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 4.8e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-52 (1-28) x AAS11714 (1-1278)

QY 1 G|Y|T|T|T|V|A|G|L|W|E|T|H|V|A|G|L|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|N|I|L|E|U|V|A|A|S|P 20
DB 136 G|G|C|T|A|C|T|A|G|T|G|A|G|T|G|A|C|C|G|G|G|C|A|C|C|C|C|C|G|C|A|G|C|G|C|T|C|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|H|G|I|S|E|R|S|E|R|A|S|P|H|E|A|I|A|V|A| 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|T|C|A|G|T|G 219

RESULT 3

AAD17877
ID AAD17877 standard; cDNA; 1278 BP.
XX

AC AAD17877;
XX

DT 10-DEC-2001 (first entry)
 XX T7-Caspase-Caspase 8 cleavage-human-pro-Asp2(a) lacking TM domain cDNA.
 DE
 XX Human; aspartyl protease 1; Aspl; amyloid precursor protein; APP;
 XX Alzheimer's disease; AD; dementia; neurofibrillary tangle; gliosis;
 KW amyloid plaque; neuronal loss; proteolytic; neurotrophic; neuroprotective;
 KW T7-Caspase-Caspase 8 cleavage-human-pro-Asp 2(a) protein; ss.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FT 1..1278
 FT /tag= a
 FT /product= "T7-Caspase-Caspase 8 cleavage-Human-pro- Asp
 FT 2(a) protein lacking transmembrane domain"
 XX
 XX GB2357767-A.
 XX
 PD 04-JUL-2001.
 XX
 PF 22-SEP-2000; 2000GB-00023315.
 XX
 PR 23-SEP-1999; 99US-00404133.
 PR 23-SEP-1999; 99US-0155493P.
 PR 23-SEP-1999; 99WO-US020881.
 PR 13-OCT-1999; 99US-00416901.
 PR 06-DEC-1999; 99US-0169232P.
 XX
 PA (PHMA) PHARMACIA & UPJOHN CO.
 PI
 PI Bienkowski MJ, Gurney M;
 DR WPI: 2001-444208/48.
 DR P-PsDB; AAE10641.
 PT Polypeptide comprising fragments of human aspartyl protease with amyloid
 PT precursor protein processing activity and alpha-secretase activity, for
 PT identifying modulators useful in treating Alzheimer's disease.
 XX
 PS Example 9; Page 128; 187pp; English.
 XX
 CC The patent discloses human aspartyl protease 1 (hu-Aspl) or modified Aspl
 CC proteins which lack transmembrane domain or amino terminal domain or
 CC cytoplasmic domain and retains alpha-secretase activity and amyloid
 CC protein precursor (APP) processing activity. The proteins of the
 CC invention are useful for assaying hu-Aspl alpha-secretase activity, which
 CC in turn is useful for identifying modulators of hu-Aspl alpha-secretase
 CC activity, where modulators that increase hu-Aspl alpha-secretase activity
 CC are useful for treating Alzheimer's disease (AD) which causes progressive
 CC dementia with consequent formation of amyloid plaques, neurofibrillary
 CC tangles, gliosis and neuronal loss. Hu-Aspl protease substrate is useful
 CC for assaying hu-Aspl proteolytic activity, by contacting hu-Aspl protein
 CC with the substrate under acidic conditions and determining the level of
 CC hu-Aspl proteolytic activity. The present sequence is a cDNA encoding T7-
 CC Caspase-Caspase 8 cleavage-human-pro-Asp 2(a) protein lacking a
 CC transmembrane (TM) domain which is generated from human Asp 2(a) protein
 CC by the addition of T7 tag and caspase 8 leader sequence at its N-terminal
 CC end and deletion of its C-terminal transmembrane domain
 XX
 SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;
 Alignment Scores:
 Pred. No.: 4,8e-14 Length: 1278
 Score: 144.00 Matches: 28
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 4 Gaps: 0
 US-10-726-967A-52 (1-28) x AAD17877 (1-1278)

QY 1 GIYTYTYValGlumetThrValGlySerProProGlnThrLeuAsnIleLeuValAsp 20
 DB 136 GGGTACTAGTGGAGATGACGCTGGGCGAGCCCCCGCAGACGCTCAACATCTGGTGAT 195
 QY 21 ThrGlySerSerAspPheAlaVal 28
 DB 196 ACAGCGAGCAGTACTTTCAGCTG 219
 RESULT 4
 AAD13033
 ID AAD13033 standard; cDNA; 1278 BP.
 XX
 XX AAD13033;
 AC
 XX 23-OCT-2001 (first entry)
 DT
 XX
 DE T7-Caspase-Caspase 8 cleavage-Human-pro-Asp2(a) deltatm protein cDNA.
 XX
 KW Human; aspartyl protease 2a; Asp 2a; beta-amyloid precursor protein; APP;
 KW beta-secretase; Alzheimer's disease; dementia; amyloid plaque; gliosis;
 KW neurofibrillary tangle; neuronal loss; amyloid-beta peptide; neurotrophic;
 KW neuroprotective; antisense therapy; gene therapy;
 KW caspase-caspase 8 cleavage-pro-Asp2(a) deltatm protein; ss.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 FT 1..1278
 FT /tag= a
 FT /product= "T7-Caspase-Caspase 8 cleavage-Human-pro-
 FT Asp2(a) deltatm protein"
 FT
 XX
 PN WO200150829-A2.
 XX
 PD 19-JUL-2001.
 XX
 PF 09-MAY-2001; 2001WO-1B000799.
 PF 09-MAY-2001; 2001WO-1B000799.
 PR
 XX
 XX (BIEN/) BIENKOWSKI M J.
 PA (GURNEY/) GURNEY M B.
 PA (HEIN/) HEINRIKSON R L.
 PA (PARO/) PARODI L A.
 PA (YANR/) YAN R.
 XX
 PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
 DR WPI: 2001-483072/52.
 DR P-PsDB; AAE06871.
 PT Novel purified polypeptide comprising fragment of mammalian aspartyl
 PT protease 2, lacking Asp2 transmembrane domain and retaining beta
 PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
 PT activity.
 XX
 PS Example 9; Page 158; 185pp; English.
 XX
 CC The invention relates to human aspartyl proteases (hu-Asp), beta-amyloid
 CC precursor protein (APP) isoforms and their corresponding DNA molecules.
 CC Human aspartyl proteases can act as beta-secretase proteases useful for
 CC treating Alzheimer's disease. APP isoforms are useful for identifying
 CC modulators of amyloid-beta peptide production, for use in designing
 CC therapeutics for the treatment and prevention of Alzheimer's disease,
 CC dementia, formation of amyloid plaques, neurofibrillary tangles, gliosis
 CC and neuronal loss. APP isoforms are also used in methods for identifying
 CC inhibitors and modulators of human Asp2 activity. The invention relates
 CC to a method for identifying agents that modulate the activity of human
 CC aspartyl protease Asp2. Amyloid-beta peptides obtained from APP are used
 CC as a means to screen in cellular assays for the inhibitors of beta- and
 CC gamma- secretase. Hu-Asp DNA fragments are useful as probes or primers in
 CC polymerase chain reactions (PCR). The probes are useful for detecting Hu-

CC Asp nucleic acids in in vitro assays and in Northern and Southern blots.
 CC The present cDNA sequence encodes T7-Caspase-Caspase 8 cleavage- Human-
 CC pro-aspartyl protease 2a (Asp2a) delta TM protein which is obtained by the
 CC addition of T7 tag and caspase 8 leader sequence at the N-terminal end
 CC and deletion of the transmembrane domain at the C-terminal end of Hu-
 CC Asp2a. Human Asp2a has beta-secretase activity

SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	4.8e-14	Length:	1278
Score:	144.00	Matches:	28
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	4	Gaps:	0

US-10-726-967a-52 (1-28) x AAD06751 (1-1278)

QY 1 G1YTYTtYtValG1uMeTtThrValG1ySerProG1nThrLeuAn1leuValAap 20

Db 136 GGCTACTACCTGAGATGACCGTGGCAGCCCGCCGACGCTCAACATCCTGTGGAT 195

QY 21 ThrG1ySerSerAenPheAlaVal 28

Db 196 ACAGGACAGCACTTAATTGCAAGT 219

RESULT 5

AAD06751 standard; cDNA; 1278 BP.

AC AAD06751;

DT 10-AUG-2001 (first entry)

DE T7-Caspase-Caspase 8-cleavage-human-pro-Asp-2(a) delta TM protein CDNA.

KM Human; alpha-secretase; amyloid precursor protein; APP; therapy;

KW Alzheimer's disease; antialzheimer's; aspartyl protease 2a; Asp2a;

OS Homo sapiens.

OS Synthetic.

Key	Location/Qualifiers
CDS	1..1278
	/*tag= a
	/product= "T7-Caspase-Caspase 8-cleavage-human-pro- Asp-2(a) delta TM protein"

FN W020013533-A2.

PD 05-APR-2001.

PF 22-SEP-2000; 2000WO-US026080.

PR 23-SEP-1999; 99US-0155493P.

PR 13-SEP-1999; 99WO-US020881.

PR 13-OCT-1999; 99US-00416901.

PR 06-DEC-1999; 99US-0169232P.

PA (PHAA) PHARMACIA & UPJOHN CO.

PI Gurney M, Bienkowski MJ;

DR WPI; 2001-290516/30.

DR P-PSDB; AAB02593.

XX Enzymes that cleave the alpha-secretase site of the amyloid precursor

PT protein, useful for the treatment of Alzheimer's disease.

XX Example 9; Page 157; 189pp; English.

CC The present invention relates to enzymes for cleaving the alpha-
 CC secretase site of the amyloid precursor protein (APP) and methods of
 CC identifying those enzymes. The methods may be used to identify enzymes
 CC that may be used to cleave the alpha-secretase cleavage site of the APP
 CC protein. The enzymes may be used to treat or modulate the progress of
 CC Alzheimer's disease. The present sequence is a cDNA encoding human
 CC Aspartyl protease 2a (Asp-2a) caspase-caspase 8-delta TM protein which is
 CC obtained by deleting the transmembrane domain and adding a T7-caspase
 CC leader sequence at the N-terminal end. This sequence has beta-secretase
 CC protease activity

SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

Alignment Scores:	4.8e-14	Length:	1278
Pred. No.:	144.00	Matches:	28
Score:	100.00%	Conservative:	0
Percent Similarity:	100.00%	Mismatches:	0
Best Local Similarity:	100.00%	Indels:	0
Query Match:	100.00%	Gaps:	0
DB:	4		

US-10-726-967a-52 (1-28) x AAD06751 (1-1278)

QY 1 G1YTYTtYtValG1uMeTtThrValG1ySerProG1nThrLeuAn1leuValAap 20

Db 136 GGCTACTACCTGAGATGACCGTGGCAGCCCGCCGACGCTCAACATCCTGTGGAT 195

QY 21 ThrG1ySerSerAenPheAlaVal 28

Db 196 ACAGGACAGCACTTAATTGCAAGT 219

RESULT 6

AAS11529 standard; cDNA; 1278 BP.

AC AAS11529;

DT 24-OCT-2001 (first entry)

DE T7-Caspase-caspase 8-Human-pro-Asp 2(a) delta TM fusion protein CDNA.

KM Human; Aspartyl protease; beta-secretase; neurotrophic; ASP2;

KW neuroprotective; amyloid protein precursor; APP; Alzheimer's disease;

KW amyloid-beta; Abeta; T7-Caspase-caspase 8-Human-pro-Asp 2(a) delta TM;

OS se.

OS Homo sapiens.

OS Synthetic.

Key	Location/Qualifiers
CDS	1..1278
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	/product= "T7-Caspase-caspase 8-Human-pro-Asp 2(a) delta TM fusion protein"
sig_peptide	43..87
	/*tag= b
	/note= "Caspase leader sequence"
mat_peptide	88..1275
	/label= Mature Asp 2(a)
	/note= "Also encodes 5 extra N-terminal amino acids constituting a caspase 8 cleavage site"

FN W0200149096-A2.

PD 12-JUL-2001.

PF 09-MAY-2001; 2001WO-IB000798.

PR 09-MAY-2001; 2001WO-IB000798.

PR 09-MAY-2001; 2001WO-IB000798.

PA (BIEN/) BIENKOWSKI M J.

PA (GURN/) GURNEY M E.

PA (HEIN/) HEINRIKSON R L.

PA (PARO/) PARODI L A.
 PA (YANR/) YAN R.
 PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
 XX MPI: 2001-502549/55.
 XX P-PSDB; AAU06615.
 DR
 XX Novel purified polypeptide comprising fragment of mammalian aspartyl
 PT protease 2, lacking Asp2 transmembrane domain and retaining beta
 PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
 PT activity.
 XX
 PS Example 9, Page 158, 185pp; English.
 XX
 CC The invention relates to a purified polypeptide comprising a fragment of
 CC mammalian aspartyl protease (Asp)2 protein which lacks the Asp2
 CC transmembrane domain and the Asp2 protein, and where the polypeptide and
 CC the fragment retain the beta-secretase activity of the mammalian Asp2
 CC protein. The invention also details polynucleotides for the Asp proteins
 CC and vectors expressing them, and a polypeptide (isoform of amyloid
 CC protein precursor (APP)) comprising the amino acid sequence of an APP or
 CC its fragment containing an APP cleavage site recognizable by a mammalian
 CC beta-secretase, and further comprising two lysine residues at the
 CC carboxyl terminus of the amino acid sequence of the mammalian APP or APP
 CC fragment. Also included in the invention are methods of identifying
 CC modulators or inhibitors of Asp2. Modulators and inhibitors of Asp2 are
 CC useful for treating Alzheimer's disease. APP is useful in methods for
 CC identifying inhibitors or modulators of human Asp2 activity and amyloid-
 CC beta (Abeta) peptide production. APP is also useful in designing
 CC therapeutics for the treatment or prevention of Alzheimer's disease. APP
 CC comprising the APP-Sw-beta-secretase peptide sequence (NLND), which is
 CC associated with increased levels of A-beta processing is useful in assays
 CC relating to the Alzheimer's research. The expression vector is useful for
 CC recombinantly expressing APP. Nucleic acids that hybridize to Asp
 CC oligonucleotides are useful as probes or primers. The probes are useful
 CC for detecting hu-Asp nucleic acids in in vitro assays and in Northern and
 CC Southern blots. The present sequence encodes T7-caspase-8 human-
 CC pro-Asp 2(a) delta TM fusion protein which has a N-terminal T7 tag to aid
 CC purification when expressed in E. coli, a Caspase leader sequence and a
 CC caspase 8 cleavage signal to aid cleavage of the signal peptide
 CC
 SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 4.8e-14 Length: 1278
 Score: 144.00 Matches: 28
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 4 Gaps: 0

US-10-726-967A-52 (1-28) x AAS11529 (1-1278)

QY 1 G|Y|T|Y|T|Y|V|A|G|U|E|C|T|H|V|A|G|Y|S|E|P|P|R|O|G|I|N|H|I|E|U|S|A|N|I|E|L|E|U|V|A|A|S|P 20
 DB 136 G|G|C|T|A|C|T|G|A|G|A|T|G|A|C|C|G|G|G|G|A|G|C|C|C|C|G|A|G|A|C|G|T|C|A|C|A|T|C|T|G|T|G|A|T 195
 QY 21 T|H|G|Y|S|E|S|E|A|N|P|H|E|A|I|A|V|A| 28
 DB 196 A|C|A|G|G|C|A|G|C|A|A|C|T|T|G|C|A|G|T|G 219

RESULT 7
 ABL52469
 ID ABL52469 standard; cDNA; 1278 BP.
 AC ABL52469;
 XX
 DT 16-JUL-2002 (first entry)
 XX
 XX T7-caspase-caspase 8 cleavage-human-pro-Asp-2(a)deltaTM nucleotide.
 DE Human; Asp-1; Asp-2; aspartyl protease; Alzheimer's disease; proteolytic;
 XX

KW amyloid precursor protein; APP; gene; ss.
 XX Homo sapiens.
 OS
 XX Key Location/Qualifiers
 FH CDS 1..1278
 FT 1..1278
 FT /tag= a
 FT /product= "T7-caspase-caspase 8 cleavage-human-pro- Asp-
 FT 2(a)deltaTM"
 XX
 PN GB2367060-A.
 XX
 PD 27-MAR-2002.
 XX
 PP 29-OCT-2001; 2001GB-00025934.
 XX
 PR 23-SEP-1999; 99US-00404133.
 PR 23-SEP-1999; 99US-0155493P.
 PR 23-SEP-1999; 99WO-US020881.
 PR 13-OCT-1999; 99US-00416901.
 PR 06-DEC-1999; 99US-0169232P.
 PR 22-SEP-2000; 2000GB-00023315.
 XX
 PA (PHAA) PHARMACIA & UPJOHN CO.
 XX
 PI Bienkowski MJ, Gurney M;
 XX
 DR MPI; 2002-397167/43.
 DR P-PSDB; ABB78602.
 XX
 PT Human aspartyl protease 1 substrates useful in assays to detect aspartyl
 PT protease activity, e.g. for the diagnosis of Alzheimer's disease.
 XX
 PS Example 9, Page 128; 182pp; English.
 XX
 CC The present invention describes a human aspartyl protease 1 (hu-Asp1)
 CC substrate (I) which comprises a peptide of no more than 50 amino acids,
 CC and which comprises the 8 amino acid sequence Gly-Leu-Ala-Leu-Ala-Leu-
 CC Glu-Pro. Also described are: (1) a method (II) for assaying hu-Asp1
 CC proteolytic activity, comprising: (a) contacting a hu-Asp1 protein with
 CC (1) under acidic conditions; and (b) determining the level of hu-Asp1
 CC proteolytic activity; (2) a purified polynucleotide (III) comprising a
 CC nucleotide sequence that hybridizes under stringent conditions to the non-
 CC coding strand complementary to a defined 1804 nucleotide sequence (see
 CC ABL52456) where the nucleotide sequence encodes a polypeptide having Asp1
 CC proteolytic activity and lacks nucleotides encoding a transmembrane
 CC domain; (3) a purified polynucleotide (III') comprising a sequence that
 CC hybridizes under stringent conditions to (III) (the nucleotide sequence
 CC encodes a polypeptide further lacking a pro-peptide domain corresponding
 CC to amino acids 23-62 of hu-Asp1 (see ABB78589); (4) a vector (IV)
 CC comprising (III) or (III'); and (5) a host cell (V) transformed or
 CC transfected with (III), (III') and/or (IV). The hu-Asp1 protease
 CC substrate (I) may be used as an enzyme substrate in assays to detect
 CC aspartyl protease activity, (II) and therefore diagnose diseases
 CC associated with aberrant hu-Asp1 expression and activity such as
 CC Alzheimer's disease. Hu-Asp1 has been localized to chromosome 21, while
 CC hu-Asp2 has been localized to chromosome 11q23.3-24.1. The present
 CC sequence encodes T7-caspase-caspase 8 cleavage-human-pro-Asp-2(a)deltaTM,
 CC which is given in an example from the present invention
 CC
 SQ Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 4.8e-14 Length: 1278
 Score: 144.00 Matches: 28
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: 6 Gaps: 0

US-10-726-967A-52 (1-28) x ABL52469 (1-1278)

QY 1 G|Y|T|Y|T|Y|V|A|G|U|E|C|T|H|V|A|G|Y|S|E|P|P|R|O|G|I|N|H|I|E|U|S|A|N|I|E|L|E|U|V|A|A|S|P 20

```

Db      136  GGCCTACTACGTGAGATGACCGTGGCGACCCCGCGACGCTCAACATCTGTGAT 195
QY      21  ThrGlySerSerAenPheAlaVal 28
Db      196  ACAGGACAGCAGTAACTTTGCAGTG 219

RESULT 8
ADJ94339 ADJ94339 standard; cDNA; 1278 BP.
XX
AC      ADJ94339;
XX
DT      03-JUN-2004 (first entry)
XX
DE      Human T7-Caspase-Caspase 8 cleavage-human-pro-Asp-2(a)deltaTM cDNA.
XX
KM      Human; sg; gene; aspartyl protease; Asp-1; Asp-2(a); Asp-2(b);
KW      beta secretase; amyloid protein precursor; APP; Alzheimer's disease;
KW      neurotrophic; neuroprotective; amyloid beta; mutant.
XX
OS      Homo sapiens.
OS      Synthetic.
XX
PN      US6706485-B1.
XX
PD      16-MAR-2004.
XX
PF      12-APR-2000; 2000US-00548376.
XX
PR      24-SEP-1998; 98US-0101594P.
PR      23-SEP-1999; 99US-00404133.
PR      23-SEP-1999; 99US-0155493P.
PR      23-SEP-1999; 99WO-US020881.
PR      13-OCT-1999; 99US-00416901.
XX
PA      (PHAA ) PHARMACIA & UPJOHN CO.
XX
PI      Gurney ME, Blenkowski MJ, Heinrikson RL, Parodi LA, Yan R;
XX
DR      WPI; 2004-236722/22.
DR      P-PSDB; ADJ94340.
XX
PT      Identifying agents that modulate activity of Asp2 aspartyl protease
PT      useful for treating or preventing Alzheimer's disease involves comparing
PT      APP processing activity of protease in presence and absence of test
PT      agent.
XX
PS      Disclosure; SEQ ID NO 27; 109pp; English.
XX
XX
CC      The invention relates to identifying agents that modulate activity of
CC      Asp2 (e.g. a beta-secretase, e.g. human Asp-2(b) appearing as ID 6,
CC      encoded by ID 5) aspartyl protease, involves contacting Asp2 with amyloid
CC      precursor protein (APP) in the presence and absence of a test agent,
CC      where Asp2 is a recombinant polypeptide and processes APP into amyloid
CC      beta, determining APP processing activity of Asp2 in presence and absence
CC      of the test agent, and comparing the activities to identify agents that
CC      modulate the activity of Asp2. Also disclosed are the cDNA and proteins
CC      for human Asp-1 and Asp-2(a), mouse Asp-2(b), a vector comprising the
CC      nucleic acid encoding Hu-Asp2 protease sequence, a host cell comprising
CC      the vector and the method of producing Hu-Asp polypeptide, an isolated
CC      antibody that specifically binds to Hu-Asp polypeptides, identifying a
CC      cell that can be used to screen for inhibitors of beta secretase
CC      activity, novel isoforms of amyloid protein precursor (APP), where the
CC      last 2 carboxy terminus amino acids of that isoform are both lysine
CC      mutation where KM at 555-596 is mutated to NU, designated e.g. APP695-Sw
CC      or APP695-Sw-KK, or a V to F mutation at 642, e.g. APP695-VF, all useful
CC      for assaying for beta secretase activity and screening for inhibitors of
CC      beta-secretase) and polynucleotides that encode the APP proteins. The
CC      method is useful for identifying agents that modulate the activity
CC      (amyloid precursor protein processing activity) of Asp2 aspartyl
CC      protease. Preferably, the method is useful for identifying agents that

```

```

CC      inhibit Asp2 aspartyl protease activity. The inhibitors of amyloid
CC      precursor protein processing, are useful for treating or preventing
CC      Alzheimer's disease. The present sequence encodes an aspartyl protease
CC      mutant construct (e.g. lacking a transmembrane domain and/or including a
CC      caspase cleavage site) used to investigate the cleavage activity of Asp2
CC      proteins.
XX
SQ      Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;
XX
Alignment Scores:
Pred. No.: 4.8e-14
Score: 144.00 Length: 1278
Percent Similarity: 100.00% Matches: 28
Best Local Similarity: 100.00% Conservative: 0
Query Match: 100.00% Mismatches: 0
DB: 12 Indels: 0 Gaps: 0
XX
US-10-726-967A-52 (1-28) x ADJ94339 (1-1278)
QY      1  G|YTYTYrValGlumetThrVal|GlySerProProGlnThrLeuAnllleuValAsp 20
Db      136  GGCCTACTACGTGAGATGACCGTGGCGACCCCGCGACGCTCAACATCTGTGAT 195
QY      21  ThrGlySerSerAenPheAlaVal 28
Db      196  ACAGGACAGCAGTAACTTTGCAGTG 219

RESULT 9
ADJ94339 ADJ94339 standard; DNA; 1278 BP.
XX
AC      ADJ94339;
XX
DT      29-JUN-2004 (first entry)
XX
DE      T7-Caspase-Caspase 8 cleavage-Human-pro-Asp-2(a)deltaTM DNA.
XX
KM      Aspartyl protease; Asp; beta secretase; amyloid precursor protein; APP;
KW      Alzheimer's disease; gene therapy; caspase; human; gene; chimeric; ds.
XX
OS      Homo sapiens.
OS      Chimeric.
OS      Unidentified.
XX
FH      Key Location/Qualifiers
FT      CDS 1..1278
FT      /cage= a
FT      /product= "T7-Caspase-Caspase 8 cleavage-Human-pro-Asp-
FT      2(a)deltaTM protein"
XX
XX
XX      US6737510-B1.
XX
XX      18-MAY-2004.
XX
PF      12-APR-2000; 2000US-00548373.
XX
PR      24-SEP-1998; 98US-0101594P.
PR      23-SEP-1999; 99US-00404133.
PR      23-SEP-1999; 99US-0155493P.
PR      23-SEP-1999; 99WO-US020881.
PR      13-OCT-1999; 99US-00416901.
XX
PA      (PHAA ) PHARMACIA & UPJOHN CO.
XX
PI      Gurney ME, Blenkowski MJ, Heinrikson RL, Parodi LA, Yan R;
XX
DR      WPI; 2004-387112/36.
DR      P-PSDB; ADJ94336.
XX
PT      New Asp2 aspartyl protease protein comprising tripeptides DTG and DSG
PT      involved in processing amyloid precursor protein into amyloid beta,
PT      useful in preparing a composition for treating or preventing Alzheimer's
PT      disease.

```

XX Example 9; SEQ ID NO 27; 108pp; English.

XX The invention relates to a method for identifying an agent that decreases

XX the protease activity of the aspartyl protease (Asp) polypeptide. It also

XX provides enzyme and enzymatic procedures for cleaving the beta secretase

XX cleavage site of the amyloid precursor protein (APP). The invention is

XX useful in preparing a composition for treating or preventing Alzheimer's

XX disease. It is also useful in gene therapy. The present sequence is T7-

XX Caspase-Caspase 8 cleavage-Human-pro-Asp-2(a)deltaTM chimeric DNA. This

XX sequence is used to illustrate the method of the invention.

XX

XX Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

XX

XX Alignment Scores:

XX Pred. No.: 4,8e-14 Length: 1278

XX Score: 144.00 Matches: 28

XX Percent Similarity: 100.00% Conservative: 0

XX Best Local Similarity: 100.00% Mismatches: 0

XX Query Match: 100.00% Indels: 0

XX DB: 12 Gaps: 0

XX

XX US-10-726-967a-52 (1-28) x ADO50435 (1-1278)

XX

XX 1 GYTYTYTYValGluMetThrValGlySerProProGlnThrLeuAsnIleuValAsp 20

XX 136 GGCTACTACGTGAGATGACCGTGGGACGCCGCCGACGCTCAACATCTGTGGAT 195

XX

XX 21 ThrGlySerSerAspAsphea1aVal 28

XX 196 ACAGGACGACGTAACCTTTCACAGT 219

XX

XX RESULT 10

XX ADR75348

XX ID ADR75348 standard; DNA; 1278 BP.

XX

XX ADR75348;

XX

XX 18-NOV-2004 (first entry)

XX

XX T7-Caspase-Caspase 8 cleavage-Human-pro-Asp-2(a)deltaTM DNA.

XX

XX Aspartyl protease; Asp; amyloid precursor protein; APP; amyloid beta;

XX Chromosome identification; Alzheimer's disease; human; caspase; chimeric;

XX gene; de.

XX

XX Homo sapiens.

XX OS Chimeric.

XX OS Unidentified.

XX

XX Key Location/Qualifiers

XX FT CDS 1..1278

XX FT /product= "T7-Caspase-Caspase 8 cleavage-Human-pro-Asp-

XX FT 2(a)deltaTM protein"

XX

XX US2004166507-A1.

XX

XX 26-AUG-2004.

XX

XX 29-AUG-2003; 2003US-00652045.

XX

XX 24-SEP-1998; 98US-0101594P.

XX 23-SEP-1999; 99US-00404133.

XX 23-SEP-1999; 99US-0155493P.

XX 13-OCT-1999; 99US-00416901.

XX

XX (GURN/) GURNEY M E.

XX (BIEN/) BIENKOWAKI M J.

XX (HEIN/) HEINRIKSON R L.

XX (PARO/) PARODI L A.

XX (YANR/) YAN R.

XX

PI Gurney ME, Bienkowiak MJ, Heinrichson RL, Parodi LA, Yan R;

XX MPI: 2004-624916/60.

XX DR P-PSDB; ADR75349.

XX

XX Novel purified/isolated polynucleotide encoding polypeptide having

XX aspartyl protease activity involved in processing amyloid precursor

XX protein into amyloid beta, useful in identifying agent decreasing

XX activity of aspartyl protease.

XX

XX Example 9; SEQ ID NO 27; 107pp; English.

XX

XX The invention relates to nucleic acid sequences encoding aspartyl

XX protease (Asp) polypeptides having aspartyl protease activity involved in

XX processing amyloid precursor protein (APP) into amyloid beta. The

XX invention also relates to a method for identifying an agent that

XX decreases the protease activity of the Asp. Asp DNA is useful in

XX chromosome identification as they can hybridise with a specific location

XX on a human chromosome and in identifying the relationship between genes

XX and diseases (particular gene responsible for causing diseases). It is

XX also useful for identifying candidates to modulate the progression of

XX Alzheimer's disease. Asp is useful in raising antibodies that are useful

XX in diagnostic assay for detecting Hu-Asp polypeptide expression. The

XX present sequence is the T7-Caspase-Caspase 8 cleavage-Human-pro-Asp-

XX 2(a)deltaTM DNA. This sequence is used to illustrate the method of the

XX invention.

XX

XX Sequence 1278 BP; 284 A; 356 C; 353 G; 285 T; 0 U; 0 Other;

XX

XX Alignment Scores:

XX Pred. No.: 4,8e-14 Length: 1278

XX Score: 144.00 Matches: 28

XX Percent Similarity: 100.00% Conservative: 0

XX Best Local Similarity: 100.00% Mismatches: 0

XX Query Match: 100.00% Indels: 0

XX DB: 13 Gaps: 0

XX

XX US-10-726-967a-52 (1-28) x ADR75348 (1-1278)

XX

XX 1 GYTYTYTYValGluMetThrValGlySerProProGlnThrLeuAsnIleuValAsp 20

XX 136 GGCTACTACGTGAGATGACCGTGGGACGCCGCCGACGCTCAACATCTGTGGAT 195

XX

XX 21 ThrGlySerSerAspAsphea1aVal 28

XX 196 ACAGGACGACGTAACCTTTCACAGT 219

XX

XX RESULT 11

XX AAD17895

XX ID AAD17895 standard; cDNA; 1287 BP.

XX

XX AAD17895;

XX

XX 10-DEC-2001 (first entry)

XX

XX Human-Asp 2(b) protein lacking transmembrane domain encoding cDNA.

XX

XX Human; aspartyl protease 2b; Asp2b; amyloid precursor protein; APP;

XX Alzheimer's disease; AD; dementia; neurofibrillary tangle; gliosis;

XX amyloid plaque; neuronal loss; proteolytic; neurotropic; neuroprotective;

XX se.

XX

XX Homo sapiens.

XX OS Synthetic.

XX

XX Key Location/Qualifiers

XX FT CDS 1..1287

XX FT /tag= a

XX FT /product= "Human-Asp 2(b) protein lacking transmembrane

XX FT domain"

XX

XX GB2357767-A.

XX

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PD   04-JUL-2001.
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PX
```

```
PF     22-SEP-2000; 2000GB-00023315.
```

```
PR
```

```
XX      23-SEP-1999;    99US-00404133.
```

```
PR       23-SEP-1999;    99US-0155493P.
```

```
PR       23-SEP-1999;    99WO-USO20881.
```

```
PR      13-OCT-1999;    99US-0046901.
```

```
PR      06-DEC-1999;    99US-0169232P.
```

```
PX
```

```
PA      (PHAA ) PHARMACIA & UPJOHN CO.
```

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PI      Bienkowski MJ, Gurney M;
```

```
DH      WPI; 2001-444208/48.
```

```
DR      P-PDB; AAEI0646.
```

```
XX
```

```
PT      Polypeptide comprising fragments of human aspartyl protease with amyloid  
PX precursor protein processing activity and alpha-secretase activity, for  
PY identifying modulators useful in treating Alzheimer's disease.  
PS  
SS Example 10; Page 137; 187pp; English.
```

```
CC The patent discloses human aspartyl protease 1 (hu-Asp1) or modified Asp1  
CC proteins which lack transmembrane domain or amino terminal domain or  
CC cytoplasmic domain and retains alpha-secretase activity and amyloid  
CC protein precursor (APP) processing activity. The proteins of the  
CC invention are useful for assaying hu-Asp1 alpha-secretase activity,  
CC in turn is useful for identifying modulators of hu-Asp1 alpha-secretase  
CC activity, where modulators that increase hu-Asp1 alpha-secretase  
CC are useful for treating Alzheimer's disease (AD) which causes progressive  
CC dementia with consequent formation of amyloid plaques, neurofibrillary  
CC tangles, gliosis and neuronal loss. Hu-Asp1 protease substrate is useful  
CC for assaying hu-Asp1 proteolytic activity, by contacting hu-Asp1 protein  
CC with the substrate under acidic conditions and determining the level of  
CC hu-Asp1 proteolytic activity. The present sequence is a cDNA encoding  
CC human Asp 2(b) protein lacking a transmembrane (TM) domain which is  
CC generated by the deletion of the C-terminal TM domain and intracellular  
CC domains of human Asp 2(b) protein  
CX  
SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;  
  
Alignment Scores:  
Pred. No.:           4,84e-14          Length:         1287  
Score:             144.00              Matches:        28  
Percent Similarity: 100.00%            Conservative:    0  
Best Local Similarity: 100.00%          Mismatches:    0  
Query Match:       100.00%             Indels:        0  
Dn:               4                   Gaps:          0  
  
US-10-726-967A-52 (1-28) x AAID17895 (1-1287)
```

```
OY      1 GlyTYYTYRVALGIMetThrvAlGIserProPGInThrleuaSnlleuvAlasp 20  
Db      220 GGCTACTAGTGAGATGCACCgGGCACAccccCGcAcAgCcTCtCAATccTGIGANT 279  
OY      21 ThrGISerSerSeAsnphealVal 28  
Db      280 ACAGCACAGTAActTTGCCAGTG 303
```

```
RESULT 12  
AADJ3276  
ID      AADI3276 standard; CDNA; 1287 BP.  
XX  
XX      AADJ3276;  
XX      AC  
DX      23-Oct-2001 (first entry)  
XX  
DD      Human-Asp2(b) deltaTM protein CDNA.  
KM      Human; asparyl protease 2b; Asp 2b; beta-amyloid precursor protein; APP;  
KW beta-secretase; Alzheimer's disease; dementia; amyloid plaque; gliosis;  
KW neurofibrillary tangle; neuronal loss; amyloid-beta peptide; nootropic;
```

KW neuroprotective; antisense therapy; Asp2(b) deltaTM protein;
KW gene therapy; ss.
OS Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
XX CDS 1..1287
XX FT /tag=a
XX FT /product= "Human Asp2(b) deltaTM protein"
XX PN
XX MO200150829-A2.
XX PD
XX 19-JUL-2001.
XX PF
XX 09-MAY-2001; 2001WO-IB000799.
XX PR
XX 09-MAY-2001; 2001WO-IB000799.
XX PA (BIEN/) BIENKOWSKI M J.
XX PA (GURN/) GURNEY M E.
XX PA (HEIN/) HEINRIKSON R L.
XX PA (PARO/) PARODI L A.
XX PA (YANK/) YAN R.
XX PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
XX DR WPI: 2001-483072/52.
XX P-PSTB; MAE06891.
XX PT Novel purified polypeptide comprising fragment of mammalian aspartyl
XX PT protease 2, lacking Asp2 transmembrane domain and retaining beta
XX PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
XX PT activity.
XX PS Example 10; Page 166-167; 185pp; English.
XX CC The invention relates to human aspartyl proteases (Hu-Asp), beta-amyloid
XX CC precursor protein (APP) isoforms and their corresponding DNA molecules.
XX CC Human aspartyl proteases can act as beta-secretase proteases useful for
XX CC treating Alzheimer's disease. APP isoforms are useful for identifying
XX CC modulators of amyloid-beta peptide production, for use in designing
XX CC therapeutics for the treatment and prevention of Alzheimer's disease,
XX CC dementia, formation of amyloid plaques, neurofibrillary tangles, gliosis
XX CC and neuronal loss. APP isoforms are also used in methods for identifying
XX CC inhibitors and modulators of human Asp2 activity. The invention relates
XX CC to a method for identifying agents that modulate the activity of human
XX CC aspartyl protease Asp2. Amyloid-beta peptides obtained from APP are used
XX CC as a means to screen in cellular assays for the inhibitors of beta- and
XX CC gamma- secretases. Hu-Asp DNA fragments are useful as probes or primers in
XX CC polymerase chain reactions (PCR). The probes are useful for detecting Hu-
XX CC Asp nucleic acids in in vitro assays and in Northern and Southern blots.
XX CC The present cDNA sequence encodes Human aspartyl protease 2b (Hu-Aspb2).
XX CC deltaTM protein which is obtained by the deletion of C-terminal
XX CC transmembrane and intracellular domains of Hu-Aspb2. Human Aspb2 has beta
XX CC -secretase activity

SQ Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:	Mismatches:	Indels:	Gaps:
Score:	4,84e-14	1287	28			
Percent Similarity:	144.00					
Best Local Similarity:	100.00%					
Query Match:	100.00%					
DB:	4					

US-10-726-967A-52 (1-28) x AAD13276 (1-1287)

GY 1 GLYYTYTValAlglMetThrValGlYSeRProFroGlnThrLeuAnlileuVal1Aap 20
db 220 GGCTATTACGTGGAGATACCGTGAGGACGCCCCCGCAGACGCTCAACATCCTGGTGGAT 279

21	ThrglySerSerAspPheAlaVal	28	
280	ACAGGACAGTACTTGCAGTG	303	
RESULT 13			
ID	AAD06768		
XX	AAD06768 standard; cDNA; 1287 BP.		
XX	AAD06768;		
XX	10-AUG-2001 (first entry)		
DE	Human aspartyl protease 2 (b) delta TM cDNA.		
KW	Human; alpha-secretase; amyloid precursor protein; APP; therapy;		
KW	Alzheimer's disease; antialzheimer's; aspartyl protease 2; Asp 2;		
XX	beta-secretase; chromosome 11q23.3-24.1; mutant; ss.		
OS	Homo sapiens.		
OS	Synthetic.		
FH	Key	Location/Qualifiers	
FT	CDS	1..1287	
FT		/*tag= a	
FT		/product= "Human aspartyl protease 2 (b) delta TM"	
XX	WO200123533-A2.		
PN			
PD	05-APR-2001.		
XX			
FP	22-SEP-2000; 2000MO-US026080.		
XX			
PR	23-SEP-1999;	99US-0155493P.	
PR	23-SEP-1999;	99MO-US020881.	
PR	13-OCT-1999;	99US-00416901.	
PR	06-DEC-1999;	99US-0169232P.	
XX			
PA	(PHAA) PHARMACIA & UPJOHN CO.		
XX			
PI	Gurney M, Bienkowski MJ;		
XX			
DR	WPI; 2001-290516/30.		
DR	P-PSDB; AAE02598.		
XX			
PT	Enzymes that cleave the alpha-secretase site of the amyloid precursor		
XX	protein, useful for the treatment of Alzheimer's disease.		
XX			
PS	Example 10; Page 165-166; 189pp; English.		
XX			
CC	The present invention relates to enzymes for cleaving the alpha-		
CC	secretase site of the amyloid precursor protein (APP) and methods of		
CC	identifying those enzymes. The methods may be used to identify enzymes		
CC	that may be used to cleave the alpha-secretase cleavage site of the APP		
CC	protein. The enzymes may be used to treat or modulate the progress of		
CC	Alzheimer's disease. The present sequence is human aspartyl protease 2		
CC	(Asp 2) (b) delta TM cDNA. The Asp 2 gene from which it is derived is		
CC	located on chromosome 11q23.3-24.1. The Asp 2 has beta-secretase protease		
CC	activity		
XX			
XX			
Sequence	1287 BP;	271 A;	370 C; 384 G; 262 T; 0 U; 0 Other;
Alignment Scores:			
Pred. No.:	4.84e-14	Length:	1287
Score:	144.00	Matches:	28
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	4	Gaps:	0
US-10-726-967A-52 (1-28) x AAD06768 (1-1287)			

Db	220	GGCTACTACGTGGAGTACCGTGGGCGACGCCCCGCGAGACGTCAACATCTGGTGGAT	27
Qy	21	ThrglySerSerAsnPhenAlaVal 28	
Db	280	ACAGCGACGAGTAACTTTGGCACTG 303	
RESULT 14			
Id	AA511547	standard; cDNA; 1287 BP.	
XX	AA511547;		
XX	24-OCT-2001	(first entry)	
XX			
DE		Human cDNA encoding Human-pro-Asp 2 (b) delta TM.	
XX			
KW		Human; Aspartyl protease; beta-secretase; neurotropic; Asp2;	
KW		neuroprotective; amyloid protein precursor; APP; Alzheimer's disease;	
KW		amyloid-beta; Abeta; Human-pro-Asp 2(b) delta TM; ss; mutant.	
XX			
OS		Homo sapiens.	
OS		Synthetic.	
XX			
FT	Key	Location/Qualifiers	
FT	CDS	1..1287	
FT		/tag= a	
FT		/product= "Human-pro-Asp 2(b) delta TM"	
XX			
FN	MO200149098-A2.		
XX			
PD	12-JUL-2001.		
XX			
XX	09-MAY-2001; 2001MO-IB000798.		
XX			
PR	09-MAY-2001; 2001MO-IB000798.		
XX			
PA	(BIEN/) BIENKOWSKI M J.		
PA	(GURN/) GURNEY M E.		
PA	(HEIN/) HEINRIKSON R L.		
PA	(PARO/) PARODI L A.		
PA	(YANR/) YAN R.		
PI	Bienskowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;		
DR	WPI; 2001-502549/55.		
XX			
PT		Novel purified polypeptide comprising fragment of mammalian aspartyl	
PT		protease 2, lacking Asp2 transmembrane domain and retaining beta	
PT		secretase activity of Asp2 useful for identifying inhibitors of Asp2	
PT		activity.	
XX			
PS		Disclosure; Page 166-167; 1855pp; English.	
XX			
CC		The invention relates to a purified polypeptide comprising a fragment of	
CC		mammalian aspartyl protease (Asp2) protein which lacks the Asp2	
CC		transmembrane domain and the Asp2 protein, and where the polypeptide and	
CC		the fragment retain the beta-secretase activity of the mammalian Asp2	
CC		protein. The invention also details polynucleotides for the Asp proteins	
CC		and vectors expressing them, and a polypeptide (isoform of amyloid	
CC		protein precursor (APP) comprising the amino acid sequence of an APP or	
CC		its fragment containing an APP cleavage site recognizable by a mammalian	
CC		beta-secretase, and further comprising two lysine residues at the	
CC		carboxyl terminus of the amino acid sequence of the mammalian APP or APP	
CC		fragment. Also included in the invention are methods of identifying	
CC		modulators or inhibitors of Asp2. Modulators and inhibitors of Asp2 are	
CC		useful for treating Alzheimer's disease. APP is useful in methods for	
CC		identifying inhibitors or modulators of human Asp2 activity and amyloid-	
CC		beta (Abeta) peptide production. APP is also useful in designing	
CC		therapeutics for the treatment or prevention of Alzheimer's disease. APP	
CC		comprising the APP-Sw-beta-secretase peptide sequence (NDA), which is	
CC		associated with increased levels of Abeta processing is useful in assays	
CC		relating the Alzheimer's research. The expression vector is useful for	
CC		recombinantly expressing APP. Nucleic acids that hybridize to Asp	

CC oligonucleotides are useful as probes or primers. The probes are useful
 CC for detecting Hu-Asp nucleic acids in in vitro assays and in Northern and
 CC Southern blots. The present sequence encodes Human-pro- Asp 2 (b) delta TM
 CC protein, which lacks the C-terminal transmembrane domain

XX Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 4,84e-14 Length: 1287
 Score: 144.00 Matches: 28
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: Gaps: 0

US-10-726-967A-52 (1-28) x ABL52487 (1-1287)

QY 1 GlyTyrTyrValGluMetThrValGlySerProProGlnThrLeuAenIleLeuValAsp 20
 Db 220 GGCTACTACGTGAGATGACCGTGGCAGCCCCCGCAGACGCTCAACATCTGTGGAT 279
 QY 21 ThrGlySerSerAenPheAlaVal 28
 Db 280 ACAGGACAGCACTTTCAGGTG 303

RESULT 15
 ABL52487
 ID ABL52487 standard; cDNA; 1287 BP.
 XX
 AC ABL52487;
 XX

DT 16-JUN-2002 (first entry)

XX Human Asp-2 (b) deltatm nucleotide sequence SEQ ID NO:50.

XX Human; Asp-1; Asp-2; aspartyl protease; Alzheimer's disease; proteolytic;
 KM Chromosome 11q23.3-24.1; gene; ss.
 XX

OS Homo sapiens.

XX Key Location/Qualifiers
 FT CDS 1..1287
 FT /tag= a
 FT /product= "Human Asp-2 (b) delta TM"

GB2367060-A.

27-MAR-2002.

PF 29-OCT-2001; 2001GB-00025934.

PR 23-SEP-1999; 99US-00404133.

PR 23-SEP-1999; 99US-015493P.

PR 23-SEP-1999; 99MO-US020881.

PR 13-OCT-1999; 99US-00416901.

PR 06-DEC-1999; 99US-0169232P.

PR 22-SEP-2000; 2000GB-00023315.

XX (PHAA) PHARMACIA & UPJOHN CO.

XX Bienkowskaki MJ, Gurney M;

XX WPI; 2002-397167/43.

XX P-PSDB; ABB78607.

XX Human aspartyl protease 1 substrates useful in assays to detect aspartyl
 PT protease activity, e.g. for the diagnosis of Alzheimer's disease.
 XX
 PS Example 10; Page 137; 182p; English.
 CC The present invention describes a human aspartyl protease 1 (hu-Asp1)
 CC substrate (I) which comprises a peptide of no more than 50 amino acids,
 CC and which comprises the 8 amino acid sequence Gly-Leu-Ala-Leu-

CC Glu-Pro. Also described are: (1) a method (II) for assaying hu-Asp1
 CC proteolytic activity, comprising: (a) contacting a hu-Asp1 protein with
 CC (I) under acidic conditions; and (b) determining the level of hu-Asp1
 CC proteolytic activity; (2) a purified polynucleotide (III) comprising a
 CC nucleotide sequence that hybridises under stringent conditions to the non
 CC -coding strand complementary to a defined 1804 nucleotide sequence (see
 CC ABL52456) where the nucleotide sequence encodes a polypeptide having Asp1
 CC proteolytic activity and lacks nucleotides encoding a transmembrane
 CC domain; (3) a purified polynucleotide (III') comprising a sequence that
 CC hybridises under stringent conditions to (III) (the nucleotide sequence
 CC encodes a polypeptide further lacking a pro-peptide domain corresponding
 CC to amino acids 23-62 of hu-Asp1 (see ABB78589)); (4) a vector (IV)
 CC comprising (III) or (III'); and (5) a host cell (V) transformed or
 CC transfected with (III), (III') and/or (IV). The hu-Asp1 protease
 CC substrate (I) may be used as an enzyme substrate in assays to detect
 CC aspartyl protease activity, (II) and therefore diagnose diseases
 CC associated with aberrant hu-Asp1 expression and activity such as
 CC Alzheimer's disease. Hu-Asp1 has been localised to chromosome 21, while
 CC hu-Asp2 has been localised to chromosome 11q23.3-24.1. The present
 CC sequence encodes human Asp-2 (b) deltatm, which is given in an example from
 CC the present invention

XX Sequence 1287 BP; 271 A; 370 C; 384 G; 262 T; 0 U; 0 Other;

Alignment Scores:
 Pred. No.: 4,84e-14 Length: 1287
 Score: 144.00 Matches: 28
 Percent Similarity: 100.00% Conservative: 0
 Best Local Similarity: 100.00% Mismatches: 0
 Query Match: 100.00% Indels: 0
 DB: Gaps: 0

US-10-726-967A-52 (1-28) x ABL52487 (1-1287)

QY 1 GlyTyrTyrValGluMetThrValGlySerProProGlnThrLeuAenIleLeuValAsp 20
 Db 220 GGCTACTACGTGAGATGACCGTGGCAGCCCCCGCAGACGCTCAACATCTGTGGAT 279
 QY 21 ThrGlySerSerAenPheAlaVal 28
 Db 280 ACAGGACAGCACTTTCAGGTG 303

Search completed: July 27, 2005, 17:25:06
 Job time : 440 secs

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 16:49:48 ; Search time 132 Seconds
(without alignments)
347.089 Million cell updates/sec

Title: US-10-726-967A-52

Perfect score: 144
Sequence: 1 GYVEMTGSPPQTINILVDTGSSNPAV 28

Scoring table:

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Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:

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-LOOPT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blomsum62 -TRANS=human40.cdi
-LIST=45 -DOCLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=pct -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US10726967@cgn2_1.1.105@runat.26072005.130735.6165 -NCPV=6 -ICPV=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV_TIMEOUT=120 -MARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

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5: /cgn2_6/prodata/1/ina/PCTUS.COMB.seq.*
6: /cgn2_6/prodata/1/ina/backfile1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	144	100.0	1278	3	US-09-548-372D-27
2	144	100.0	1278	3	US-09-548-367D-27
3	144	100.0	1278	4	US-09-551-853D-27
4	144	100.0	1278	4	US-09-548-376D-27
5	144	100.0	1278	4	US-09-548-376D-27
6	144	100.0	1278	4	US-09-548-376D-27
7	144	100.0	1278	4	US-09-548-373D-27
8	144	100.0	1278	4	US-09-548-373D-27
9	144	100.0	1278	4	US-09-548-366F-27
10	144	100.0	1278	4	US-09-548-366F-27
11	144	100.0	1278	4	US-09-548-368D-27
12	144	100.0	1278	4	US-09-794-925A-27

13	144	100.0	1278	4	US-09-806-194A-27	Sequence 27, Appl
14	144	100.0	1287	3	US-09-548-372D-50	Sequence 50, Appl
15	144	100.0	1287	3	US-09-548-367D-50	Sequence 50, Appl
16	144	100.0	1287	4	US-09-551-853D-50	Sequence 50, Appl
17	144	100.0	1287	4	US-09-416-901B-50	Sequence 50, Appl
18	144	100.0	1287	4	US-09-548-376D-50	Sequence 50, Appl
19	144	100.0	1287	4	US-09-794-927A-50	Sequence 50, Appl
20	144	100.0	1287	4	US-09-548-373D-50	Sequence 50, Appl
21	144	100.0	1287	4	US-09-795-847B-50	Sequence 50, Appl
22	144	100.0	1287	4	US-09-869-414-50	Sequence 50, Appl
23	144	100.0	1287	4	US-09-548-366F-50	Sequence 50, Appl
24	144	100.0	1287	4	US-09-548-368D-50	Sequence 50, Appl
25	144	100.0	1287	4	US-09-794-925A-50	Sequence 50, Appl
26	144	100.0	1302	3	US-09-548-372D-25	Sequence 25, Appl
27	144	100.0	1302	3	US-09-548-367D-25	Sequence 25, Appl
28	144	100.0	1302	4	US-09-551-853D-25	Sequence 25, Appl
29	144	100.0	1302	4	US-09-416-901B-25	Sequence 25, Appl
30	144	100.0	1302	4	US-09-548-376D-25	Sequence 25, Appl
31	144	100.0	1302	4	US-09-794-927A-25	Sequence 25, Appl
32	144	100.0	1302	4	US-09-548-373D-25	Sequence 25, Appl
33	144	100.0	1302	4	US-09-795-847B-25	Sequence 25, Appl
34	144	100.0	1302	4	US-09-869-414-25	Sequence 25, Appl
35	144	100.0	1302	4	US-09-548-366F-25	Sequence 25, Appl
36	144	100.0	1302	4	US-09-548-368D-25	Sequence 25, Appl
37	144	100.0	1302	4	US-09-794-925A-25	Sequence 25, Appl
38	144	100.0	1302	4	US-09-806-194A-25	Sequence 25, Appl
39	144	100.0	1305	3	US-09-548-372D-52	Sequence 52, Appl
40	144	100.0	1305	3	US-09-548-367D-52	Sequence 52, Appl
41	144	100.0	1305	4	US-09-551-853D-52	Sequence 52, Appl
42	144	100.0	1305	4	US-09-416-901B-52	Sequence 52, Appl
43	144	100.0	1305	4	US-09-548-376D-52	Sequence 52, Appl
44	144	100.0	1305	4	US-09-794-927A-52	Sequence 52, Appl
45	144	100.0	1305	4	US-09-548-373D-52	Sequence 52, Appl

ALIGNMENTS

RESULT 1
US-09-548-372D-27
Sequence 27, Application US/09548372D
Patent No. 6420534
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
FILE OR INVENTION: THEREOF
FILE REFERENCE: 29915/62801
CURRENT APPLICATION NUMBER: US/09/548, 372D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 27
LENGTH: 1278
TYPE: DNA
ORGANISM: Homo sapiens
US-09-548-372D-27
Alignment Scores:
Pred. No.: 4.05e-16
Score: 144.00
Percent Similarity: 100.00%
Best Local Similarity: 100.00%
Query Match: 100.00%
Matches: 28
Conservative: 0
Mismatch: 0
Indels: 0
Gaps: 0
US-10-726-967A-52 (1-28) x US-09-548-372D-27 (1-1278)

QY 1 GlyTyrValGluMetThrValGlySerProProGlnThrLeuAsnIleLeuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGAGCCCCCGGAGAGCTCAACATCTGTGGAT 195
QY 21 ThrGlySerSerAspPheAlaVal 28
Db 196 ACAGGACAGCACTTAATTGACAGTG 219

RESULT 2
US-09-548-367D-27
; Sequence 27, Application US/09548367D
; Patent No. 6440698
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280H
; CURRENT APPLICATION NUMBER: US/09/548,367D
; CURRENT FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-548-367D-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-548-367D-27 (1-1278)

QY 1 GlyTyrValGluMetThrValGlySerProProGlnThrLeuAsnIleLeuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGAGCCCCCGGAGAGCTCAACATCTGTGGAT 195
QY 21 ThrGlySerSerAspPheAlaVal 28
Db 196 ACAGGACAGCACTTAATTGACAGTG 219

RESULT 3
US-09-551-853D-27
; Sequence 27, Application US/09551853D
; Patent No. 6500667
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280L
; CURRENT APPLICATION NUMBER: US/09/551,853D
; CURRENT FILING DATE: 2000-04-18
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24

; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-551-853D-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-551-853D-27 (1-1278)

QY 1 GlyTyrValGluMetThrValGlySerProProGlnThrLeuAsnIleLeuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGAGCCCCCGGAGAGCTCAACATCTGTGGAT 195
QY 21 ThrGlySerSerAspPheAlaVal 28
Db 196 ACAGGACAGCACTTAATTGACAGTG 219

RESULT 4
US-09-416-901B-27
; Sequence 27, Application US/09416901B
; Patent No. 6699671
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280A
; CURRENT APPLICATION NUMBER: US/09/416,901B
; CURRENT FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; NUMBER OF SEQ ID NOS: 72
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-416-901B-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-416-901B-27 (1-1278)

QY 1 GlyTyrValGluMetThrValGlySerProProGlnThrLeuAsnIleLeuValAsp 20
Db 136 GGCTACTACGTGAGATGACCGTGGGAGCCCCCGGAGAGCTCAACATCTGTGGAT 195
QY 21 ThrGlySerSerAspPheAlaVal 28
Db 196 ACAGGACAGCACTTAATTGACAGTG 219

RESULT 5
US-09-548-376D-27

Sequence 27, Application US/09548373D
Patent No. 6706485
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR
TITLE OF INVENTION: AND USES
TITLE OF INVENTION: THEREOF
FILE REFERENCE: 29915/6280F
CURRENT APPLICATION NUMBER: US/09/548,376D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 27
LENGTH: 1278
TYPE: DNA
ORGANISM: Homo sapiens
US-09-548-376D-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-548-376D-27 (1-1278)

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DB 136 GGCCTACTAGCTGAGTAGACCGTGGGCAAGCCCCCGACAGCGCTCAACATCTCTGGTGAT 195

QY 21 T|H|G|Y|S|E|R|S|E|R|A|S|N|P|H|E|A|I|A|V|I 28
DB 196 ACAGCGACGACGTACTTTCGACGTG 219

RESULT 6
US-09-794-927A-27
Sequence 27, Application US/09794927A
Patent No. 6727074
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
TITLE OF INVENTION: THEREOF
FILE REFERENCE: 29915/6280FG
CURRENT APPLICATION NUMBER: US/09/794,927A
CURRENT FILING DATE: 2001-02-27
PRIOR APPLICATION NUMBER: 09/416,901
PRIOR FILING DATE: 1999-10-13
PRIOR APPLICATION NUMBER: 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 74
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 27
LENGTH: 1278
TYPE: DNA
ORGANISM: Homo sapiens
US-09-794-927A-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-794-927A-27 (1-1278)

QY 1 G|Y|T|T|Y|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|T|H|R|E|U|S|N|I|E|L|E|U|V|A|A|P 20
DB 136 GGCCTACTAGCTGAGTAGACCGTGGGCAAGCCCCCGACAGCGCTCAACATCTCTGGTGAT 195

QY 21 T|H|G|Y|S|E|R|S|E|R|A|S|N|P|H|E|A|I|A|V|I 28
DB 196 ACAGCGACGACGTACTTTCGACGTG 219

RESULT 7
US-09-548-373D-27
Sequence 27, Application US/09548373D
Patent No. 6737510

GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
TITLE OF INVENTION: THEREOF
FILE REFERENCE: 29915/6280B
CURRENT APPLICATION NUMBER: US/09/548,373D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20881
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 27
LENGTH: 1278
TYPE: DNA
ORGANISM: Homo sapiens
US-09-548-373D-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-548-373D-27 (1-1278)

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DB 136 GGCCTACTAGCTGAGTAGACCGTGGGCAAGCCCCCGACAGCGCTCAACATCTCTGGTGAT 195

QY 21 T|H|G|Y|S|E|R|S|E|R|A|S|N|P|H|E|A|I|A|V|I 28
DB 196 ACAGCGACGACGTACTTTCGACGTG 219

RESULT 8
US-09-795-847B-27
Sequence 27, Application US/09795847B
Patent No. 6753163
GENERAL INFORMATION:
APPLICANT: GURNEY, MARK E.
APPLICANT: BIENKOWSKI, MICHAEL J.
APPLICANT: HEINRIKSON, ROBERT L.
APPLICANT: PARODI, LUIS A.
APPLICANT: YAN, RIGDANG

;; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
;; FILE REFERENCE: 28341/6280DE
;; CURRENT APPLICATION NUMBER: US/09/795,847B
;; PRIOR FILING DATE: 2001-02-28
;; PRIOR APPLICATION NUMBER: 09/416,901
;; PRIOR FILING DATE: 1999-10-13
;; PRIOR APPLICATION NUMBER: 60/155,493
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: 09/404,133
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: PCT/US99/20881
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: 60/101,594
;; NUMBER OF SEQ ID NOS: 74
;; SOFTWARE: PatentIn Ver. 2.0
;; SEQ ID NO 27
;; LENGTH: 1278
;; TYPE: DNA
;; ORGANISM: Homo sapiens
US-09-795-847B-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

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Db 136 GGCCTACTAGTGAATGATGACCGTGGCAGCCCCCGCAGACCTCAACATCTCTGTGAT 195
21 ThnglSerSerSnpheAlaVal 28
Db 196 ACAGGACGACGTAACCTTTCAGTG 219

RESULT 9
US-09-869-414-27
;; Sequence 27, Application US/09869414
;; Patent No. 6790610
;; GENERAL INFORMATION:
;; APPLICANT: BEINKOWSKI ET AL.
;; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
;; FILE REFERENCE: 28341/6280M
;; CURRENT APPLICATION NUMBER: US/09/869,414
;; PRIOR FILING DATE: 2001-06-27
;; PRIOR APPLICATION NUMBER: 09/416,901
;; PRIOR FILING DATE: 1999-10-13
;; PRIOR APPLICATION NUMBER: 60/155,493
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: 09/404,133
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: PCT/US99/20881
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: 60/101,594
;; NUMBER OF SEQ ID NOS: 73
;; SOFTWARE: PatentIn Ver. 2.0
;; SEQ ID NO 27
;; LENGTH: 1278
;; TYPE: DNA
;; ORGANISM: Homo sapiens
US-09-869-414-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

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Db 136 GGCCTACTAGTGAATGATGACCGTGGCAGCCCCCGCAGACCTCAACATCTCTGTGAT 195
21 ThnglSerSerSnpheAlaVal 28
Db 196 ACAGGACGACGTAACCTTTCAGTG 219

RESULT 10
US-09-548-366F-27
;; Sequence 27, Application US/09548366F
;; Patent No. 6797487
;; GENERAL INFORMATION:
;; APPLICANT: GURNEY ET AL.
;; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
;; FILE REFERENCE: 29915/6280J
;; CURRENT APPLICATION NUMBER: US/09/548,366F
;; PRIOR FILING DATE: 2000-04-12
;; PRIOR APPLICATION NUMBER: US 60/155,493
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: US 09/404,133
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: PCT/US99/20881
;; PRIOR FILING DATE: 1999-09-23
;; PRIOR APPLICATION NUMBER: US 60/101,594
;; NUMBER OF SEQ ID NOS: 73
;; SOFTWARE: PatentIn version 3.1
;; SEQ ID NO 27
;; LENGTH: 1278
;; TYPE: DNA
;; ORGANISM: Homo sapiens
US-09-548-366F-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-548-366F-27 (1-1278)
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Db 136 GGCCTACTAGTGAATGATGACCGTGGCAGCCCCCGCAGACCTCAACATCTCTGTGAT 195
21 ThnglSerSerSnpheAlaVal 28
Db 196 ACAGGACGACGTAACCTTTCAGTG 219

RESULT 11
US-09-548-368D-27
;; Sequence 27, Application US/09548368D
;; Patent No. 6825023
;; GENERAL INFORMATION:
;; APPLICANT: GURNEY ET AL.
;; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
;; FILE REFERENCE: 29915/6280C
;; CURRENT APPLICATION NUMBER: US/09/548,368D
;; PRIOR FILING DATE: 2000-04-12
;; PRIOR APPLICATION NUMBER: US 60/155,493
;; PRIOR FILING DATE: 1999-09-23

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28

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; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-548-368D-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-548-368D-27 (1-1278)
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Db 136 GGCTACTAGCTGAGATGACCGTGGCAGACCCCGCAGACGCTCAACATCTCTGTGGAT 195

QY 21 ThnGlySerSerAsnPhaAlaVal 28
Db 196 ACAGGACGACGTAACCTTTCGACGTG 219

RESULT 12
US-09-794-925A-27
; Sequence 27, Application US/09794925A
; Patent No. 6828117
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280H1
; CURRENT APPLICATION NUMBER: US/09/794,925A
; CURRENT FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-794-925A-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-794-925A-27 (1-1278)
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Db 136 GGCTACTAGCTGAGATGACCGTGGCAGACCCCGCAGACGCTCAACATCTCTGTGGAT 195
QY 21 ThnGlySerSerAsnPhaAlaVal 28
Db 196 ACAGGACGACGTAACCTTTCGACGTG 219

RESULT 13
US-09-806-194A-27
; Sequence 27, Application US/09806194A
; Patent No. 6835565
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowsk, Michael J.
; APPLICANT: Heinrikson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; APPLICANT: Pharmacia & Upjohn Company
; TITLE OF INVENTION: Alzheimer's Disease Secretase
; FILE REFERENCE: 6177.P CP
; CURRENT APPLICATION NUMBER: US/09/806,194A
; CURRENT FILING DATE: 2001-09-17
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-806-194A-27

Alignment Scores:
Pred. No.: 4,05e-16 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 4 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-806-194A-27 (1-1278)
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Db 136 GGCTACTAGCTGAGATGACCGTGGCAGACCCCGCAGACGCTCAACATCTCTGTGGAT 195

QY 21 ThnGlySerSerAsnPhaAlaVal 28
Db 196 ACAGGACGACGTAACCTTTCGACGTG 219

RESULT 14
US-09-548-372D-50
; Sequence 50, Application US/09548372D
; Patent No. 6420534
; GENERAL INFORMATION:
; APPLICANT: GURNEY ET AL.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
; FILE REFERENCE: 29915/6280I
; CURRENT APPLICATION NUMBER: US/09/548,372D
; CURRENT FILING DATE: 2000-04-12
; PRIOR APPLICATION NUMBER: US 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: US 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
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ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-548-372D-50

Alignment Scores:
Pred. No.: 4.09e-16 Length: 1287
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-548-372D-50 (1-1287)

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QY 21 ThrGlySerSerAsnPheAlaVal 28
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RESULT 15

US-09-548-367D-50
Sequence 50, Application US/09548367D
Patent No. 6440698
GENERAL INFORMATION:
APPLICANT: GURNEY ET AL.
TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR AND USES
FILE OF INVENTION: THEREOF
FILE REFERENCE: 29915/6280H
CURRENT APPLICATION NUMBER: US/09/548,367D
CURRENT FILING DATE: 2000-04-12
PRIOR APPLICATION NUMBER: US 60/155,493
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 09/404,133
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: PCT/US99/20861
PRIOR FILING DATE: 1999-09-23
PRIOR APPLICATION NUMBER: US 60/101,594
PRIOR FILING DATE: 1998-09-24
NUMBER OF SEQ ID NOS: 73
SOFTWARE: PatentIn version 3.1
SEQ ID NO 50
LENGTH: 1287
TYPE: DNA
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: Hu-Asp2 (b) delta TM
US-09-548-367D-50

Alignment Scores:

Pred. No.: 4.09e-16 Length: 1287
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-548-367D-50 (1-1287)

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QY 21 ThrGlySerSerAsnPheAlaVal 28
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Db 280 ACAGGACGAGTAACTTTGCAGTG 303
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Search completed: July 27, 2005, 18:52:41
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GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus.p2n model

Run on: July 27, 2005, 17:18:01 ; Search time 627 Seconds
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Title: US-10-726-967A-52
Perfect score: 144
Sequence: 1 GYVEMTVGSPPTQTLNIVDQSSNFAV 28

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Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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and is derived by analysis of the total score distribution.

SUMMARIES

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5	144	100.0	1278	9	US-09-794-925-27	Sequence 27, Appl
6	144	100.0	1278	9	US-09-681-442-27	Sequence 27, Appl
7	144	100.0	1278	10	US-09-681-442-27	Sequence 27, Appl
8	144	100.0	1278	10	US-09-548-366-27	Sequence 27, Appl
9	144	100.0	1278	18	US-10-652-927-27	Sequence 27, Appl
10	144	100.0	1278	18	US-10-652-830-27	Sequence 27, Appl
11	144	100.0	1278	19	US-10-652-045-27	Sequence 27, Appl
12	144	100.0	1278	20	US-10-476-935-27	Sequence 27, Appl
13	144	100.0	1278	21	US-10-940-867-27	Sequence 27, Appl
14	144	100.0	1278	21	US-10-477-076-27	Sequence 27, Appl
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16	144	100.0	1287	9	US-09-795-847-50	Sequence 50, Appl
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19	144	100.0	1287	9	US-09-794-925-50	Sequence 50, Appl
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32	144	100.0	1302	9	US-09-794-748-25	Sequence 25, Appl
33	144	100.0	1302	9	US-09-794-925-25	Sequence 25, Appl
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36	144	100.0	1302	10	US-09-548-366-25	Sequence 25, Appl
37	144	100.0	1302	18	US-10-652-927-25	Sequence 25, Appl
38	144	100.0	1302	18	US-10-652-830-25	Sequence 25, Appl
39	144	100.0	1302	19	US-10-652-045-25	Sequence 25, Appl
40	144	100.0	1302	20	US-10-476-935-25	Sequence 25, Appl
41	144	100.0	1302	21	US-10-940-867-25	Sequence 25, Appl
42	144	100.0	1305	9	US-09-794-927-52	Sequence 52, Appl
43	144	100.0	1305	9	US-09-795-847-52	Sequence 52, Appl
44	144	100.0	1305	9	US-09-794-743-52	Sequence 52, Appl
45	144	100.0	1305	9	US-09-794-748-52	Sequence 52, Appl

ALIGNMENTS

RESULT 1
US-09-794-927-27
; Sequence 27, Application US/09794927
; Patent No. US20010016324A1
GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, App Substrates Therefor, And
; TITLE OF INVENTION: US28341/6280FG
; FILE REFERENCE: 28341/6280FG
; CURRENT APPLICATION NUMBER: US/09/794, 927
; CURRENT FILING DATE: 2001-02-27

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/ PRIOR APPLICATION NUMBER: 09/416,901
/ PRIOR FILING DATE: 1999-10-13
/ PRIOR APPLICATION NUMBER: 60/155,493
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: 09/404,133
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: PCT/US99/20881
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: 60/101,594
/ PRIOR FILING DATE: 1998-09-24
/ NUMBER OF SEQ ID NOS: 73
/ SOFTWARE: Patentln Ver. 2.0
/ SEQ ID NO 27
/ LENGTH: 1278
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-794-927-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-794-927-27 (1-1278)

QY 1 G|Y|T|T|Y|V|A|G|U|e|t|h|r|V|a|G|Y|S|e|r|P|r|o|G|I|n|T|h|r|e|u|a|n|l|e|u|V|a|A|a|p 20
    |||
Db 136 G|G|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|C|G|C|A|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|h|r|G|Y|S|e|r|S|e|r|a|n|P|h|e|a|l|a|V|a|l 28
    |||
Db 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|G|C|A|G|T|G 219

RESULT 2
US-09-795-847-27
/ Sequence 27, Application US/09795847
/ Patent No. US20010018208A1
/ GENERAL INFORMATION:
/ APPLICANT: Gurney, Mark E.
/ APPLICANT: Bienkowski, Michael J.
/ APPLICANT: Heinrichson, Robert L.
/ APPLICANT: Parodi, Luis A.
/ APPLICANT: Yan, Riqiang
/ TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
/ TITLE OF INVENTION: USES
/ FILE REFERENCE: 28341/6280DE
/ CURRENT APPLICATION NUMBER: US/09/795,847
/ CURRENT FILING DATE: 2001-02-28
/ PRIOR APPLICATION NUMBER: 09/416,901
/ PRIOR FILING DATE: 1999-10-13
/ PRIOR APPLICATION NUMBER: 60/155,493
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: 09/404,133
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: PCT/US99/20881
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: 60/101,594
/ PRIOR FILING DATE: 1998-09-24
/ NUMBER OF SEQ ID NOS: 73
/ SOFTWARE: Patentln Ver. 2.0
/ SEQ ID NO 27
/ LENGTH: 1278
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-795-847-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0
```

```
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-795-847-27 (1-1278)

QY 1 G|Y|T|T|Y|V|A|G|U|e|t|h|r|V|a|G|Y|S|e|r|P|r|o|G|I|n|T|h|r|e|u|a|n|l|e|u|V|a|A|a|p 20
    |||
Db 136 G|G|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|C|G|C|A|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|h|r|G|Y|S|e|r|S|e|r|a|n|P|h|e|a|l|a|V|a|l 28
    |||
Db 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|G|C|A|G|T|G 219

RESULT 3
US-09-794-743-27
/ Sequence 27, Application US/09794743
/ Patent No. US20010021391A1
/ GENERAL INFORMATION:
/ APPLICANT: Gurney, Mark E.
/ APPLICANT: Bienkowski, Michael J.
/ APPLICANT: Heinrichson, Robert L.
/ APPLICANT: Parodi, Luis A.
/ APPLICANT: Yan, Riqiang
/ TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
/ TITLE OF INVENTION: USES
/ FILE REFERENCE: 28341/6280BC
/ CURRENT APPLICATION NUMBER: US/09/794,743
/ CURRENT FILING DATE: 2001-02-27
/ PRIOR APPLICATION NUMBER: 09/416,901
/ PRIOR FILING DATE: 1999-10-13
/ PRIOR APPLICATION NUMBER: 60/155,493
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: 09/404,133
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: PCT/US99/20881
/ PRIOR FILING DATE: 1999-09-23
/ PRIOR APPLICATION NUMBER: 60/101,594
/ PRIOR FILING DATE: 1998-09-24
/ NUMBER OF SEQ ID NOS: 73
/ SOFTWARE: Patentln Ver. 2.0
/ SEQ ID NO 27
/ LENGTH: 1278
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-09-794-743-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-794-743-27 (1-1278)

QY 1 G|Y|T|T|Y|V|A|G|U|e|t|h|r|V|a|G|Y|S|e|r|P|r|o|G|I|n|T|h|r|e|u|a|n|l|e|u|V|a|A|a|p 20
    |||
Db 136 G|G|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|C|G|C|A|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|h|r|G|Y|S|e|r|S|e|r|a|n|P|h|e|a|l|a|V|a|l 28
    |||
Db 196 A|C|A|G|G|C|A|G|C|A|G|T|A|C|T|T|G|C|A|G|T|G 219

RESULT 4
US-09-794-748-27
/ Sequence 27, Application US/09794748
/ Patent No. US20020037315A1
/ GENERAL INFORMATION:
/ APPLICANT: Gurney, Mark E.
```

```

: SEQ ID NO 27
: LENGTH: 1278
: TYPE: DNA
: ORGANISM: Homo sapiens
US-09-794-925-27

Alignment Scores:
Pred. No.: 2.82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-794-925-27 (1-1278)

QY 1 GLYTYYTVAIGLMEThThVAIGLYSerProProGInThThLeuAsnIleuValaap 20
Db GGCTACTACGCGAGATGACCGTGGCGAGCCCCCGCAGAGCTCAACATCCTGGTGGAT 195
21 ThrGlySerSerAsnPhaIVal 28
Db 196 ACAGCGACGAGTAACTTGCAGTG 219

RESULT 6
US-09-681-442-27
: Sequence 27, Application US/09681442
: Patent No. US20020081634A1
: GENERAL INFORMATION:
: APPLICANT: Guirney, Mark E.
: APPLICANT: Bienkowski, Michael J.
: APPLICANT: Heinrikson, Robert L.
: APPLICANT: Parodi, Luis A.
: APPLICANT: Yan, Riqiang
: TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
: FILE REFERENCE: 28341/6280FG
: CURRENT APPLICATION NUMBER: US/09/681,442
: PRIOR FILING DATE: 2001-04-05
: PRIOR APPLICATION NUMBER: 09/416,901
: PRIOR FILING DATE: 1999-10-13
: PRIOR APPLICATION NUMBER: 60/155,493
: PRIOR FILING DATE: 1999-09-23
: PRIOR APPLICATION NUMBER: 09/404,133
: PRIOR FILING DATE: 1999-09-23
: PRIOR APPLICATION NUMBER: PCT/US99/20881
: PRIOR FILING DATE: 1999-09-23
: PRIOR APPLICATION NUMBER: 60/101,594
: PRIOR FILING DATE: 1998-09-24
: NUMBER OF SEQ ID NOS: 73
: SOFTWARE: PatentIn Ver. 2.0
: SEQ ID NO 27
: LENGTH: 1278
: TYPE: DNA
: ORGANISM: Homo sapiens
US-09-681-442-27

Alignment Scores:
Pred. No.: 2.82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-681-442-27 (1-1278)

QY 1 GLYTYYTVAIGLMEThThVAIGLYSerProProGInThThLeuAsnIleuValaap 20
Db 136 GGCTACTACGCGAGATGACCGTGGCGAGCCCCCGCAGAGCTCAACATCCTGGTGGAT 195
21 ThrGlySerSerAsnPhaIVal 28

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DB 196 ACAGCAGCAGTACTTGCAGTG 219

RESULT 7

US-09-869-414-27

Sequence 27, Application US/09869414

Publication No. US20030077226A1

GENERAL INFORMATION:

APPLICANT: Beinikowski et al.

TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES

FILE REFERENCE: 28341/6280M

CURRENT APPLICATION NUMBER: US/09/869,414

PRIOR FILING DATE: 2001-06-27

PRIOR APPLICATION NUMBER: 09/416,901

PRIOR FILING DATE: 1999-10-13

PRIOR APPLICATION NUMBER: 60/155,493

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: 09/404,133

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: PCT/US99/20881

PRIOR FILING DATE: 1999-09-23

NUMBER OF SEQ ID NOS: 60/101,594

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 27

LENGTH: 1278

TYPE: DNA

ORGANISM: Homo sapiens

US-09-869-414-27

Alignment Scores:

Pred. No.: 2,82e-14 Length: 1278

Score: 144.00 Matches: 28

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 100.00% Indels: 0

DB: 10 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-869-414-27 (1-1278)

QY 1 G|Y|T|T|V|A|G|U|e|T|h|V|A|G|Y|S|e|P|P|o|G|I|n|T|h|l|e|u|A|n|i|l|e|u|V|a|l|A|s|p 20

DB 136 G|G|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|C|C|C|C|C|C|C|G|A|G|C|C|T|C|A|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|h|g|Y|S|e|S|e|A|n|P|h|e|A|l|A|V|a|l 28

DB 196 ACAGCAGCAGTACTTGCAGTG 219

RESULT 8

US-09-548-366-27

Sequence 27, Application US/09548366

Publication No. US20030104365A1

GENERAL INFORMATION:

APPLICANT: Gurney, Mark E.

APPLICANT: Bienikowski, Michael J.

APPLICANT: Heinrikson, Robert L.

APPLICANT: Parodi, Luis A.

APPLICANT: Yan, Riqiang

TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND

FILE REFERENCE: 28341/6280A

CURRENT APPLICATION NUMBER: US/09/548,366

PRIOR FILING DATE: 2000-04-12

PRIOR APPLICATION NUMBER: 60/155,493

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: 09/404,133

PRIOR FILING DATE: 1999-09-23

PRIOR APPLICATION NUMBER: PCT/US99/20881

PRIOR FILING DATE: 1999-09-23

NUMBER OF SEQ ID NOS: 60/101,594

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 27

LENGTH: 1278

TYPE: DNA

ORGANISM: Homo sapiens

US-10-726-967a-52 (1-28) x US-09-548-366-27 (1-1278)

QY 1 G|Y|T|T|V|A|G|U|e|T|h|V|A|G|Y|S|e|P|P|o|G|I|n|T|h|l|e|u|A|n|i|l|e|u|V|a|l|A|s|p 20

DB 136 G|G|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|C|C|C|C|C|C|C|G|A|G|C|C|T|C|A|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|h|g|Y|S|e|S|e|A|n|P|h|e|A|l|A|V|a|l 28

DB 196 ACAGCAGCAGTACTTGCAGTG 219

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 27

LENGTH: 1278

TYPE: DNA

ORGANISM: Homo sapiens

US-09-548-366-27

Alignment Scores:

Pred. No.: 2,82e-14 Length: 1278

Score: 144.00 Matches: 28

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 100.00% Indels: 0

DB: 10 Gaps: 0

US-10-726-967a-52 (1-28) x US-09-548-366-27 (1-1278)

QY 1 G|Y|T|T|V|A|G|U|e|T|h|V|A|G|Y|S|e|P|P|o|G|I|n|T|h|l|e|u|A|n|i|l|e|u|V|a|l|A|s|p 20

DB 136 G|G|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|G|G|C|A|C|C|C|C|C|C|C|G|A|G|C|C|T|C|A|A|C|T|C|T|G|T|G|A|T 195

QY 21 T|h|g|Y|S|e|S|e|A|n|P|h|e|A|l|A|V|a|l 28

DB 196 ACAGCAGCAGTACTTGCAGTG 219

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RESULT 10
US-10-652-830-27
; Sequence 27, Application US/10652830
; Publication No. US20040048303A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280N1
; CURRENT APPLICATION NUMBER: US/10/652,830
; PRIOR FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: 09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-652-830-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 18 Gaps: 0

US-10-726-967a-52 (1-28) x US-10-652-830-27 (1-1278)

QY 1 G|Y|T|Y|T|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|A|N|I|L|E|U|V|A|A|S|P 20
DB 136 G|G|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|G|G|G|C|A|G|C|C|C|C|G|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195
QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|V|A|I 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|T|G|C|A|G|T|G 219

RESULT 11
US-10-652-045-27
; Sequence 27, Application US/10652045
; Publication No. US20040166507A1
; GENERAL INFORMATION:
; APPLICANT: Gurney et al.
; TITLE OF INVENTION: Alzheimer's Disease Secretase, APP Substrates Therefor and Uses
; FILE REFERENCE: 29915/6280N2
; CURRENT APPLICATION NUMBER: US/10/652,045
; PRIOR FILING DATE: 2003-08-29
; PRIOR APPLICATION NUMBER: 09/794,925
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1998-09-23
; NUMBER OF SEQ ID NOS: 60/101,594
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-476-935-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 20 Gaps: 0

US-10-726-967a-52 (1-28) x US-10-476-935-27 (1-1278)

QY 1 G|Y|T|Y|T|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|A|N|I|L|E|U|V|A|A|S|P 20
DB 136 G|G|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|G|G|G|C|A|G|C|C|C|C|G|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195
QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|V|A|I 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|T|G|C|A|G|T|G 219
```

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; NUMBER OF SEQ ID NOS: 74
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-652-045-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 19 Gaps: 0

US-10-726-967a-52 (1-28) x US-10-652-045-27 (1-1278)

QY 1 G|Y|T|Y|T|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|A|N|I|L|E|U|V|A|A|S|P 20
DB 136 G|G|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|G|G|G|C|A|G|C|C|C|C|G|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195
QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|V|A|I 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|T|G|C|A|G|T|G 219

RESULT 12
US-10-476-935-27
; Sequence 27, Application US/10476935
; Publication No. US20040234976A1
; GENERAL INFORMATION:
; APPLICANT: Belinkowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M1
; CURRENT APPLICATION NUMBER: US/10/476,935
; PRIOR FILING DATE: 2003-11-06
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-476-935-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 20 Gaps: 0

US-10-726-967a-52 (1-28) x US-10-476-935-27 (1-1278)

QY 1 G|Y|T|Y|T|V|A|G|U|W|E|T|H|V|A|G|I|S|E|R|P|R|O|G|I|N|T|H|L|E|U|S|A|N|I|L|E|U|V|A|A|S|P 20
DB 136 G|G|T|A|C|T|A|C|G|T|G|A|G|A|T|G|A|C|C|G|G|G|G|C|A|G|C|C|C|C|G|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|T 195
QY 21 T|H|G|I|S|E|R|S|E|R|A|N|P|H|E|A|I|V|A|I 28
DB 196 A|C|A|G|G|C|A|G|C|A|G|T|T|G|C|A|G|T|G 219
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RESULT 13
US-10-940-867-27
; Sequence 27, Application US/10940867
; Publication No. US20050026256A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; APPLICANT: Pharmacia & Upjohn Company
; TITLE OF INVENTION: Alzheimer's Disease Secretase
; FILE REFERENCE: 6177.PCPA
; CURRENT APPLICATION NUMBER: US/10/940, 867
; PRIOR FILING DATE: 2004-09-14
; PRIOR APPLICATION NUMBER: US 09/806,194
; PRIOR FILING DATE: 2001-03-26
; PRIOR APPLICATION NUMBER: US 60/101,594
; NUMBER OF SEQ ID NOS: 49
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-940-867-27

Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 21 Gaps: 0

US-10-726-967A-52 (1-28) x US-10-940-867-27 (1-1278)
QY 1 GATYTYTVVAlGUmEtThrValGIySeRProPGInThrLeuAnIlleuValAap 20
Db 136 GGCTACTAGTGAGATATACCGTGGCAGCCCCCGCAGACTCAACATCTGTGGAT 195
QY 21 ThrGIySeRSeRAsnPheAlaVal 28
Db 196 ACAGGCAGCAGTAATTGTCAGTG 219

RESULT 14
US-10-477-076-27
; Sequence 27, Application US/10477076
; Publication No. US20050080232A1
; GENERAL INFORMATION:
; APPLICANT: Beinowski et al.
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND USES
; FILE REFERENCE: 28341/6280M2
; CURRENT APPLICATION NUMBER: US/10/477, 076
; PRIOR FILING DATE: 2003-11-06
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 27
; LENGTH: 1278
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-477-076-27
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Alignment Scores:
Pred. No.: 2,82e-14 Length: 1278
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 21 Gaps: 0

US-10-726-967A-52 (1-28) x US-10-477-076-27 (1-1278)
QY 1 GATYTYTVVAlGUmEtThrValGIySeRProPGInThrLeuAnIlleuValAap 20
Db 136 GGCTACTAGTGAGATATACCGTGGCAGCCCCCGCAGACTCAACATCTGTGGAT 195
QY 21 ThrGIySeRSeRAsnPheAlaVal 28
Db 196 ACAGGCAGCAGTAATTGTCAGTG 219

RESULT 15
US-09-794-927-50
; Sequence 50, Application US/09794927
; Patent No. US20010016324A1
; GENERAL INFORMATION:
; APPLICANT: Gurney, Mark E.
; APPLICANT: Bienkowski, Michael J.
; APPLICANT: Heinrichson, Robert L.
; APPLICANT: Parodi, Luis A.
; APPLICANT: Yan, Riqiang
; TITLE OF INVENTION: ALZHEIMER'S DISEASE SECRETASE, APP SUBSTRATES THEREFOR, AND
; TITLE OF INVENTION: USES
; FILE REFERENCE: 28341/6280RG
; CURRENT APPLICATION NUMBER: US/09/794,927
; PRIOR FILING DATE: 2001-02-27
; PRIOR APPLICATION NUMBER: 09/416,901
; PRIOR FILING DATE: 1999-10-13
; PRIOR APPLICATION NUMBER: 60/155,493
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 09/404,133
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: PCT/US99/20881
; PRIOR FILING DATE: 1999-09-23
; PRIOR APPLICATION NUMBER: 60/101,594
; PRIOR FILING DATE: 1998-09-24
; NUMBER OF SEQ ID NOS: 73
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 50
; LENGTH: 1287
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Hu-Aap2 (b)
US-09-794-927-50

Alignment Scores:
Pred. No.: 2,84e-14 Length: 1287
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967A-52 (1-28) x US-09-794-927-50 (1-1287)
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Db 220 GGCTACTAGTGAGATATACCGTGGCAGCCCCCGCAGACTCAACATCTGTGGAT 279
QY 21 ThrGIySeRSeRAsnPheAlaVal 28
Db 280 ACAGGCAGCAGTAATTGTCAGTG 303
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Search completed: July 27, 2005, 19:03:20
Job time : 629 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: July 27, 2005, 13:56:30 ; Search time 3149 Seconds
(without alignments)
338.457 Million cell updates/sec

Title: US-10-726-967A-52

Perfect score: 144
Sequence: 1 GYVEMTVGSPPTQINILVDITGSSNPAV 28

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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-DB=EST -OPMT=fastlap -SUFFIX=p2n.rst -MINMATCH=0.1 -LOOPEXT=0 -LIST=45
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LST=45
-DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=PCO -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
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-NO MAP -LARGEOUTER -NEG_SCORES=0 -WAIT -DSPBLLOC=100 -LONGLOG
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

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2: gb_est2:*
3: gb_hic:*
4: gb_est3:*
5: gb_est4:*
6: gb_est5:*
7: gb_est6:*
8: gb_gss1:*
9: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	144	100.0	346	5	BY103030	BY103030 BY103030
2	144	100.0	365	5	BY080676	BY080676 BY080676
3	144	100.0	1123	5	BX376891	BX376891 BX376891
4	144	100.0	1506	9	AY417360	AY417360 Homo sapi
5	144	100.0	1506	9	AY417362	AY417362 Mus muscu
6	144	100.0	3634	3	AK041285	AK041285 Mus muscu
7	144	100.0	3805	3	AK082230	AK082230 Mus muscu
8	144	100.0	3859	3	AK014464	AK014464 Mus muscu
9	144	100.0	3877	3	AK033112	AK033112 Mus muscu

10	144	100.0	3880	3	AK080498	AK080498 Mus muscu
11	144	100.0	4048	3	AK082317	AK082317 Mus muscu
12	144	100.0	4101	3	AK046175	AK046175 Mus muscu
13	143	99.3	4466	3	AK049626	AK049626 Mus muscu
14	138	95.8	461	1	AL700831	AL700831 DKFZp6861
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16	137	95.8	813	7	CN224123	CN224123 WLA053E12
17	137	95.1	458	1	AL700814	AL700814 DKFZp6861
18	130	80.3	611	7	CN484125	CN484125 hw42d08.Y
19	128	80.9	727	6	CA749486	CA749486 UT-M-FY0-
20	125	86.8	1001	5	BU128383	BU128383 603113984
21	124	86.1	761	7	CN064631	CN064631 Ag2_p8_p2
22	124	86.1	763	7	CN064511	CN064511 Ag2_p7_p9
23	124	86.1	817	7	CK139305	CK139305 AGENCOURT
24	124	86.1	819	7	CK143759	CK143759 AGENCOURT
25	124	86.1	873	6	CD755522	CD755522 AGENCOURT
26	124	86.1	893	6	CA475966	CA475966 AGENCOURT
27	124	86.1	898	6	CD757678	CD757678 AGENCOURT
28	124	86.1	915	6	CD756050	CD756050 AGENCOURT
29	122	84.7	880	5	BQ733989	BQ733989 AGENCOURT
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31	120	83.3	670	6	CA375995	CA375995 654231 NC
32	119	82.6	208	1	AI290317	AI290317 cm02D05.X
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34	119	82.6	614	2	AW153854	AW153854 I126d02.Y
35	119	82.6	648	5	BM957312	BM957312 EY76E05.Y
36	119	82.6	705	5	BP434294	BP434294 BP434294
37	119	82.6	712	7	CK681409	CK681409 ZF101-P00
38	119	82.6	751	7	CV480976	CV480976 AGENCOURT
39	119	82.6	765	4	BM006442	BM006442 603615166
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41	119	82.6	783	6	CB962018	CB962018 AGENCOURT
42	119	82.6	795	6	CB998213	CB998213 AGENCOURT
43	119	82.6	814	6	CB998866	CB998866 AGENCOURT
44	119	82.6	847	7	CK871931	CK871931 AGENCOURT
45	119	82.6	847	7	CN021417	CN021417 AGENCOURT

ALIGNMENTS

RESULT 1
BY103030
LOCUS
DEFINITION
BY103030
ACCESSION
BY103030
VERSION
BY103030.1
KEYWORDS
SOURCE
ORGANISM
Mus musculus (house mouse)
EST.
Mus musculus
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 346)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S., Nishikido, I., Osato, N., Saito, R., Suzuki, H., Yamazaki, A., Kiyosawa, H., Vagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schombach, C., Cojocari, T., Balderelli, R., Hill, D.P., Bait, C., Hume, D.A., Quackenbush, J., Schriml, L.M., Kanpin, A., Matsuda, H., Batalov, S., Beisel, K.W., Blake, J.A., Brad, D., Brusic, V., Choctha, C., Corbani, L.E., Cousins, S., Dalla, E., Dragan, T.A., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, J.J., Jarvis, E.D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lenhard, B., Lyons, P.A., Maglocz, D.R., Maltais, L., Marchionni, L., McKenzle, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Pertea, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M., Sandelin, A., Schneider, C., Sempke, C.A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Verrardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L.G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I.,

Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Kono, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Aikawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E.S., Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
22354683
12466851
COMMENT

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Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Saitama-cho, Tsukuba, Ibaraki, 305-3858, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gs.c.riken.jp, URL: http://genome.gsc.riken.jp/
Aizawa, K., Akimura, T., Aikawa, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Kono, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Waki, K., Watanabe, A., Muramatsu, M. and Hayashizaki, Y. Direct Submission
Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system-384-format sequencing pipeline with 384 multichannel sequencer. Genome Res. 10 (11), 1757-1771 (2000)
Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in Riken Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.
Please visit our web site (http://genome.gsc.riken.go.jp) for further details.

FEATURES
source
1.346
Location/Qualifiers
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ORIGIN
Alignment Scores:
Pred. No.:

1.7e-13 Length: 346

Score: 144.00
Percent Similarity: 100.00%
Best Local Similarity: 100.00%
Query Match: 100.00%
DB: 5
Gaps: 0

US-10-726-967a-52 (1-28) x BY103030 (1-346)

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QY 21 ThrGlySerSerAsnPheAlaVal 28

Db 146 ACGGCACTAGTAACCTTGCAGTG 169

RESULT 2
BY080676 365 bp mRNA linear EST 07-DEC-2002
LOCUS
DERIVATION
BY080676 RIKEN full-length enriched, 16 days embryo whole body Mus

ACCESSION
BY080676
VERSION
BY080676.1 GI:26191219
KEYWORDS
EST.

SOURCE
Mus musculus (house mouse)

ORGANISM

REFERENCE

AUTHORS

1 (bases 1 to 365)
Okazaki, Y., Furuno, M., Kasukawa, T., Aachi, J., Bono, H., Kondo, S., Nakai, I., Otsu, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schombach, C., Gajbordi, T., Balderelli, L., Hill, D.P., Bull, C., Hume, D.A., Quackenbush, J., Schriml, L.M., Kanapin, A., Matsuda, H., Batalov, S., Beisel, K.W., Blake, J.A., Bradt, D., Bruscia, V., Chothia, K., Cordani, L.E., Cousins, S., Dalla, E., Dragani, T.A., Fletcher, C.F., Forrest, A., Frazer, K.S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimond, S., Gurinovich, S., Hirokawa, N., Jackson, I.J., Jerns, E.D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedierski, R.M., King, B.L., Konagaya, A., Kurochkin, I.V., Lee, Y., Lennard, B., Lyons, P.A., Maglott, D.R., Maltas, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W.J., Perce, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J.U., Qi, D., Ramchandran, S., Ravasi, T., Reed, J.C., Reed, D.J., Reid, J., Ring, B.Z., Ringwald, M., Sandelin, A., Schneider, C., Semp, C.A., Setu, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M.S., Teasdale, R.D., Tomita, M., Verardo, R., Wagner, L., Wainwright, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L.G., Wyszewski, A., Yanagisawa, M., Yang, I., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Kono, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Aikawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E.S., Rogers, J., Birney, E. and Hayashizaki, Y.

TITLE

Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

Nature 420, 563-573 (2002)
22354683
12466851

JOURNAL
MEDLINE
PUBMED

COMMENT
CONTACT: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Saitama-cho, Tsukuba, Ibaraki, 305-3858, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gs.c.riken.jp, URL: http://genome.gsc.riken.jp/
Aizawa, K., Akimura, T., Aikawa, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Kono, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K.,

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whole body"

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Alignment Scores:

US-10-726-967A-52 (1-28) x BY080676 (1-365)

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DEFINITION	BX376891 Homo sapiens NEUROBLASTOMA COT 50-NORMALIZED Homo sapiens				
	CDNA clone CS0DD007YP18 5-PRIME, mRNA sequence.				

ACCESSION	BX376891	
VERSION	BX376891.2	GI:46556538
KEYWORDS	EST.	
SOURCE	Homo sapiens	(human)
ORGANISM	Homo sapiens	

REVIEWER 1 (Pages 1 to 1123)
AUTHORS L.I. W. B., Gruber, C., Jesse, J. and Poljans, D.
TITLE Full-length cDNA libraries and normalization
JOURNAL Unpublished (2001)
COMMENT On May 8, 2003 this sequence version replaced gi:30434929

Genoscope - Centre National de Séquençage
2 rue Gaston Crémieux, CP 5706 - 91057 Evry cedex - FRANCE
Email: segref@genoscope.cns.fr, Web : www.genoscope.cns.fr
1st strand cDNA was primed with a NotI-oligo(dT) primer. Five primed

For more information about this cluster, see <http://www.genoscope.cns.fr/cdnas-CSD000707D09Q1&c=5902.r>.

Source

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/clone_lib="Homo sapiens NEUROBLASTOMA COT 50-NORMALIZED"
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sites of the pCMSPORT 6 vector. Library was normalized."

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Alignment Scores

US-10-726-967A-52 (1-28) X BX376891 (1-1123)

Qy	Db	Qy	Db
1	657	21	717
GLTTLTYValGluwecThrValGlySerProProlnTrnLeuAsn1LeuValAsp	GGCTACTACGTGGAGATGACCGTGGGACAGCCCCCGACGAGACGCTCAACATCTCGTGGAT	ThrGlySerSerAspPheIaVal	ACAGGACAGCAGTACTCTTGCAGGTG

RESULT	4
AY417360	
LOCUS	AY417360 1506 bp DNA linear GSS 17-DEC-2001
DEFINITION	Homo sapiens BACE gene, VIRUTAL TRANSCRIPT, partial sequence,
	genomic survey sequence.
ACCESSION	AY417360
VERSION	AY417360.1 GI:39773320
KEYWORDS	GSS.

ORGANISM Homo sapiens
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 1506)
REFERENCE
AUTHORS Clark, A.G., Glanowski, S., Nelson, R., Thomas, P., Kejaival, A.,

TITLE
Inferring nonneutral evolution from human-chimp-mouse orthologous

JOURNAL
PUMED
REFERENCE
AUTHORS

Science 302 (5652), 1960-1963 (2003)
14671302
2 (bases 1 to 1506)
Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejalivel, A.,
Todd, M.A., Tanenbaum, D.M., Civallo, D.R., Lu, F., Murphy, B.,
Ferrera, S., Wang, G., Zheng, X.H., White, T.J., Sinsky, J.J.,
Adams, M.D. and Caregill, M.

TITLE	Direct Submission
JOURNAL	Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive, Rockville, MD 20850, USA
COMMENT	This sequence was made by sequencing genomic exons and ordering them based on alignment.
FEATURES	Location/Qualifiers
source	1. 1506 /organism="Homo sapiens"

/mol_type="genomic DNA"
/db_xref="taxon:9606"
<1..>1506
/gene="BACE"
/locus_tag="HCM6198"

ORIGIN

Alignment Scores:

Pred. No.: 1.01e-12 Length: 1506
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967a-52 (1-28) x AY417360 (1-1506)

QY 1 G|Y|Y|T|Y|V|A|G|U|M|E|T|H|V|A|G|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|E|N|I|L|E|U|V|A|A|S|P 20

DB 220 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|C|C|A|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|C 279

QY 21 T|H|G|Y|S|E|R|S|E|R|A|E|N|P|H|E|A|L|A|V|A| 28
DB 280 A|C|G|G|C|A|G|A|G|T|A|C|T|T|G|C|A|G|T|G 303

RESULT 5
AY417362 1506 bp DNA linear GSS 17-DEC-2003
LOCUS AY417362 Mus musculus BACE gene, VIRUAL TRANSCRIPT, partial sequence,
DEFINITION AY417362 genomic survey sequence.
ACCESSION AY417362
VERSION AY417362.1 GI:39773322
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
TITLE 1 (bases 1 to 1506)
JOURNAL Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
PUBMED Ferreria, S., Wang, G., Zheng, X.H., White, T.J., Snijsky, J.J.,
14671302 Adams, M.D. and Cargill, M.
TITLE Inferring nonneutral evolution from human-chimp-mouse orthologous
JOURNAL Science 302 (5652), 1960-1963 (2003)

REFERENCE
AUTHORS 2 (bases 1 to 1506)
JOURNAL Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
PUBMED Ferreria, S., Wang, G., Zheng, X.H., White, T.J., Snijsky, J.J.,
14671302 Adams, M.D. and Cargill, M.
TITLE Direct Submission
JOURNAL Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
PUBMED Rockville, MD 20850, USA
TITLE This sequence was made by sequencing genomic exons and ordering
COMMENT them based on alignment.

FEATURES

source
1..1506
/organism="Mus musculus"
/mol_type="genomic DNA"
/db_xref="taxon:10090"
<1..>1506
/gene="BACE"
/locus_tag="HCM6198"

ORIGIN

Alignment Scores:

Pred. No.: 1.01e-12 Length: 1506
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0

US-10-726-967a-52 (1-28) x AY417362 (1-1506)

QY 1 G|Y|Y|T|Y|V|A|G|U|M|E|T|H|V|A|G|Y|S|E|R|P|R|O|G|I|N|T|H|L|E|U|A|E|N|I|L|E|U|V|A|A|S|P 20
DB 220 G|G|C|T|A|C|T|A|G|T|G|A|G|A|T|G|A|C|C|G|T|G|G|C|A|G|C|C|C|C|C|C|A|G|C|T|C|A|C|A|C|T|C|T|G|T|G|A|C 279

QY 21 T|H|G|Y|S|E|R|S|E|R|A|E|N|P|H|E|A|L|A|V|A| 28
DB 280 A|C|G|G|C|A|G|A|G|T|A|C|T|T|G|C|A|G|T|G 303

RESULT 5
AK041285 3634 bp mRNA linear HTC 03-APR-2004
LOCUS AK041285 Mus musculus adult male aorta and vein cDNA, RIKEN full-length
DEFINITION enriched library, clone:A530097B07 product:beta-site APP cleaving
enzyme, full insert sequence.
ACCESSION AK041285
VERSION AK041285.1 GI:26334342
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
JOURNAL Carninci, P. and Hayashizaki, Y.
PUBMED High-efficiency full-length cDNA cloning
99279253 Meth. Enzymol. 303, 19-44 (1999)
REFERENCE 10349636

REFERENCE
AUTHORS 2
TITLE Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
JOURNAL Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
PUBMED Normalization and subtraction of cap-trapper-selected cDNAs to
20499374 prepare full-length cDNA libraries for rapid discovery of new genes
Genome Res. 10 (10), 1617-1630 (2000)
REFERENCE 11042159

REFERENCE
AUTHORS 3
TITLE Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
JOURNAL Kono, H., Akiyama, J., Nishi, K., Katsunai, T., Tashiro, H., Itoh, M.,
PUBMED Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
20530913 Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kasahigaki, K.,
Fujisawa, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watanabe, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multichipillary sequencer
Genome Res. 10 (11), 1757-1771 (2000)
REFERENCE 11076861

REFERENCE
AUTHORS 4
TITLE The RIKEN Genome Exploration Research Group Phase II Team and the
JOURNAL FANTOM Consortium.
PUBMED Functional annotation of a full-length mouse cDNA collection
Nature 409, 685-690 (2001)
REFERENCE 5

REFERENCE
AUTHORS 5
TITLE The FANTOM Consortium and the RIKEN Genome Exploration Research
JOURNAL Group Phase I & II Team.
PUBMED Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
REFERENCE 6 (bases 1 to 3634)

REFERENCE
AUTHORS Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T.,
Horii, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Kato, H., Kawai, J., Kojima, Y., Kondo, S., Kono, H., Kouda, M.,
Koyama, S., Kurihara, C., Matsumura, T., Miyazaki, A., Murata, M.,
Nakamura, M., Nishi, K., Nomura, K., Numasaki, R., Ohno, M., Ohnishi, N.,
Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
Sano, H., Sasaki, D., Shibata, K., Shingawa, A., Shiraki, T.,
Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,

URL: <http://fantom.gsc.riken.jp/>.
Location/Qualifiers

FEATURES
source
1..3805
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="FANTOM_DB:C230026008"
/db_xref="taxon:10090"
/clone="C230026008"
/tissue_type="cerebellum"
/clone_lib="RIKEN full-length enriched mouse cDNA library"
/dev_stage="0 day neonate"
48..1935
/note="beta-site APP cleaving enzyme (MGD|MG1:1346542, GB|NM_011792, evidence: BLASTN, 98%, match=3874) putative"

misc_feature

ORIGIN

Alignment Scores:

Pred. No.:	3,11e-12	Length:	3805
Score:	144.00	Matches:	28
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	3	Gaps:	0

US-10-726-967a-52 (1-28) x AK082230 (1-3805)

QY 1 G1YTYTVA1G1uWctThVa1G1SerProProGlnrThrLeuValaap 20
667 GGCTACTATGTGGAGNAGCCGTAGGACCCCTCCACAGCGCTCAACATCTGTGTGAC 726

QY 21 ThG1YserSeranphealaval 28
727 ACGGCGAGTAGTAACTTTCGACGTG 750

Db

RESULT 8
AK014464
LOCUS
DEFINITION
Mus musculus 16 days embryo head cDNA, RIKEN full-length enriched library, clone:4122401C04 product:beta-site APP cleaving enzyme, full insert sequence.
ACCESSION
AK014464
VERSION
AK014464.1 GI:12852334
KEYWORDS
HTC; CAP trapper.
SOURCE
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS
Carninci, P. and Hayashizaki, Y.
TITLE
High-efficiency full-length cDNA cloning
JOURNAL
Meth. Enzymol. 303, 19-44 (1999)
MEDLINE
99279253
PUBMED
10349636

REFERENCE
AUTHORS
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
TITLE
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL
Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE
20499374
PUBMED
11042159

REFERENCE
AUTHORS
Shibata, K., Itoh, M., Aizawa, K., Nagao, S., Sasaki, N., Carninci, P., Kono, H., Akiyama, Y., Nishi, K., Kitsu, T., Tashiro, H., Itoh, M., Suni, N., Ishii, Y., Nakamura, S., Hazama, M., Nishino, T., Hazada, A., Yamamoto, R., Matsunoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsumura, S., Kawai, U., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE
RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer

JOURNAL
MEDLINE
PUBMED
REFERENCE
AUTHORS
TITLE
JOURNAL
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS

Genome Res. 10 (11), 1757-1771 (2000)
20530913
11076861

4
The RIKEN Genome Exploration Research Group Phase II Team and the PANTOM Consortium.
Functional annotation of a full-length mouse cDNA collection
Nature 409, 685-690 (2001)

5
The PANTOM Consortium and the RIKEN Genome Exploration Research Group Phase I & II Team.
Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)
6 (bases 1 to 3859)

Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H., Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y., Furuno, M., Hanagaki, T., Hara, A., Hayatsu, N., Hiramoto, K., Hirakawa, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Izawa, M., Kaekawa, T., Kato, H., Kawai, U., Kojima, Y., Kono, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C., Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D., Shibata, K., Shibata, Y., Shingawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Tanaka, T., Tejima, Y., Toya, T., Yamamura, T., Yasunishi, A., Yoshida, K., Yoshino, M., Muramatsu, M. and Hayashizaki, Y.
Direct Submission

Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail: genome-res@gs.c.riken.jp, URL: <http://genome.gsc.riken.jp/>, Tel: 81-45-503-9222, Fax: 81-45-503-9216)

Please visit our web site (<http://genome.gsc.riken.jp/>) for further details.

CNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. First strand cDNA was primed with a primer [5' GAGAGAGAGAGAGATCCAGAGCTCTTTTCTTTTCTTTVN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGAGATTCGATTAATTAAATTAATCCCTCCCTCC 3']. cDNA was cleaved with BamHI and XhoI. cDNA of size compressed longer than 7 kb was selected before cloning. Vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I. Cloning sites, 5' end: SalI; 3' end: BamHI. Host: DH10B.

Location/Qualifiers

1..3859
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="FANTOM_DB:4122401C04"
/db_xref="taxon:10090"
/clone="4122401C04"
/tissue_type="head"
/clone_lib="RIKEN full-length enriched mouse cDNA library"
/dev_stage="16 days embryo"
430..1935
/note="unnamed protein product; beta-site APP cleaving enzyme (MGD|MG1:1346542, GB|NM_011792, evidence: BLASTN, 98%, match=3874) putative"
/codon_start=1
/protein_id="BAB29370.1"
/db_xref="GI:12852335"
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AAITSEDFKPFNGSNWEGILGLAYAEIARPDLSLPPFDLVKQTHIPNIFSLQCGA
GFLNQTALASVGSMTIGGIDHSIYTSGLTSPYRREMYEIVIRVEIGDOLKIM
DCKEYNNKSIYDSGTTNRLPKVPEAAVKSITKASKEPDPGFWGLVCMQAG
TTPKNIFFVLSILYLMGEVTNQSFRITILPQOILRPVEDVATSDDCYKRAVQSSTGT
VNGAVIMEGFYVDFRARRRIGFAVSACHVDEFPRAAEGFPVTADMDGCGNIPOT
DESTLMTIAYVMAAICALFMLPLCLMVCQWRCLRCLRHQHDPEADISILK"

ORIGIN

Alignment Scores:

Pred. No.:	3,16e-12	Length:	3859
Score:	144.00	Matches:	28
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	3	Gaps:	0

US-10-726-967a-52 (1-28) x AK014464 (1-3859)

Qy 1 GIVTYTYVValGluMetThrValGlySerProGlnThrLeuAniLeuValAap 20

Db 649 GGCCTACTATGTGAGATGACCTGAGCAGCCCCCACAACGCTCAACCTCTGCTGAGC 708

Qy 21 ThrGlySerSerAspPheAlaVal 28

Db 709 ACGGCGAGTACTGACTTTCAGTGTG 732

RESULT 9 AK033112 3877 bp mRNA linear HTC 03-APR-2004

LOCUS AK033112 Mus musculus 15 days embryo male testis cDNA, RIKEN full-length
DEFINITION enriched library, clone:8030431G04 product:beta-site APP cleaving
enzyme, full insert sequence.

ACCESSION AK033112 GI:26328834

VERSION AK033112.1

KEYWORDS HTC; CAP trapper.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE

1 Carninci, P. and Hayashizaki, Y.
High-efficiency full-length cDNA cloning
Meth. Enzymol. 303, 19-44 (1999)

JOURNAL 99279253

PUBMED 10349636

REFERENCE

2 Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
Genome Res. 10 (10), 1617-1630 (2000)

JOURNAL 20499374

PUBMED 11042159

REFERENCE

3 Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Kum, H., Akiyama, J., Nishi, K., Katsunari, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watanabe, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsumura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system-384-format
sequencing pipeline with 384 multicapillary sequencer
Genome Res. 10 (11), 1757-1771 (2000)

JOURNAL

PUBMED

11076861

TITLE

The RIKEN Genome Exploration Research Group Phase II Team and the
FANTOM Consortium.
Functional annotation of a full-length mouse cDNA collection
Nature 409, 685-690 (2001)

JOURNAL

PUBMED

11076861

TITLE

PUBMED

11076861

TITLE

PUBMED

11076861

TITLE

PUBMED

11076861

TITLE

PUBMED

11076861

Group Phase I & II Team.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
Nature 420, 563-573 (2002)

JOURNAL

PUBMED

11076861

TITLE

PUBMED

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PUBMED

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11076861

Qy 1 GlyTyrValGluMetThrValGlySerProGlnThrLeuAsnIleLeuValAsp 20
Db 669 GCCTACTATGTGGAGATACCGTACGACGCCCCACAGACGCTCAACATCTGTGGAC 728
Qy 21 ThrGlySerSerAspHeaIaVal 28
Db 729 ACGGCGAGTACTGACTTGCAGTG 752

RESULT 10
AK080498 3880 bp mRNA linear HTC 03-APR-2004
LOCUS AK080498
DEFINITION Mus musculus 7 days neonate cerebellum cDNA, RIKEN full-length
enriched library, clone:A730059K08 product:beta-site APP cleaving
enzyme, full insert sequence.
ACCESSION AK080498
VERSION AK080498.1 GI:26099278
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Carninci, P. and Hayashizaki, Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636

AUTHORS 2
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Komno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
JOURNAL Prepare full-length cDNA libraries for rapid discovery of new genes
MEDLINE Genome Res. 10 (10), 1617-1630 (2000)
PUBMED 20499374

AUTHORS 3
Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Komno, H., Akiyama, J., Nishi, K., Katsunai, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Ikegami, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Nishine, T., Kashiwagi, K.,
Fujisake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watanabe, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsubara, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kita, A. and Hayashizaki, Y.
TITLE RIKEN integrated sequence analysis (RISA) system-384-format
JOURNAL sequencing pipeline with 384 multichipillary sequencer
MEDLINE Genome Res. 10 (11), 1757-1771 (2000)
PUBMED 20530913

AUTHORS 4
The RIKEN Genome Exploration Research Group Phase II Team and the
PANTOM Consortium.
TITLE Functional annotation of a full-length mouse cDNA collection
JOURNAL Nature 409, 685-690 (2001)
MEDLINE 11076861

AUTHORS 5
The FANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE Analysis of the mouse transcriptome based on functional annotation
JOURNAL of 60,770 full-length cDNAs
MEDLINE Nature 420, 563-573 (2002)
PUBMED 6 (baaes 1 to 3880)

REFERENCE
AUTHORS Adachi, J., Aizawa, K., Akimura, T., Arikawa, T., Bono, H., Carninci, P.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T.,
Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Kasch, H., Kawai, J., Kojima, Y., Kondo, S., Komno, H., Kouda, M.,
Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,
Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ono, M., Ohsato, N.,
Okazaki, Y., Saito, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S.,
Takeda, Y., Tanaka, T., Tomaru, A., Toyota, T., Yasunishi, A.,
Muramatsu, M. and Hayashizaki, Y.

TITLE Direct Submission
JOURNAL Submitted (16-APR-2002) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail:genome-res@gscc.riken.jp,
URL:ftp://genome.gsc.riken.jp/, Tel:81-45-503-9222,
Fax:81-45-503-9216)
COMMENT CDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse cDNAs.
Please visit our web site for further details.
URL:ftp://genome.gsc.riken.jp/
URL:ftp://fantom.gsc.riken.jp/
URL:ftp://location.qualifiers

FEATURES
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ORIGIN
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Pred. No.: 3,18e-12 Length: 3880
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Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0

US-10-726-967a-52 (1-28) x AK080498 (1-3880)

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RESULT 11
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LOCUS AK082317
DEFINITION Mus musculus 0 day neonate cerebellum cDNA, RIKEN full-length
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enzyme, full insert sequence.
ACCESSION AK082317
VERSION AK082317.1 GI:26349644
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Carninci, P. and Hayashizaki, Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636

AUTHORS 2
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Komno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.

TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
JOURNAL prepare full-length cDNA libraries for rapid discovery of new genes
MEDLINE Genome Res. 10 (10), 1617-1630 (2000)
PUBMED 20499374
REFERENCES 11042159
AUTHORS 3
Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Komano, H., Akiyama, J., Nishi, K., Kitesuna, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsunura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE RIKEN integrated sequence analysis (RISA) system-384-format
JOURNAL RIKEN integrated sequence analysis (RISA) system-384-format
MEDLINE Genome Res. 10 (11), 1757-1771 (2000)
PUBMED 20530913
REFERENCES 11076861
AUTHORS 4
The RIKEN Genome Exploration Research Group Phase II Team and the
PANTOM Consortium.
TITLE Functional annotation of a full-length mouse cDNA collection
JOURNAL Nature 409, 685-690 (2001)
MEDLINE Nature 409, 685-690 (2001)
PUBMED 11076861
AUTHORS 5
The PANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE Analysis of the mouse transcriptome based on functional annotation
JOURNAL of 60,770 full-length cDNAs
MEDLINE Nature 420, 563-573 (2002)
PUBMED 11076861
AUTHORS 6 (bases 1 to 4048)
Nature 420, 563-573 (2002)
TITLE Direct Submission
JOURNAL Submitted (16-Apr-2002) Yoshihide Hayashizaki, The Institute of
MEDLINE Physical and Chemical Research (RIKEN), Laboratory for Genome
PUBMED Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp,
URL: http://genome.gsc.riken.jp/, Tel: 81-45-503-9222,
Fax: 81-45-503-9216)
COMMENT cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site for further details.
URL: http://genome.gsc.riken.jp/
URL: http://fantom.gsc.riken.jp/
FEATURES
source location/Qualifiers
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ORIGIN
Alignment Scores:
Pred. No.: 3,356-12 Length: 4048
Score: 144.00 Matches: 28
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 3 Gaps: 0
US-10-726-967a-52 (1-28) x AK082317 (1-4048)
Qy 1 GLVYTYTYValGluMetThrValGlySerProGlnThrLeuAsnIleLeuValAsp 20
Db 670 GGCTTACTATGTGAGATGACCGTGGCAGCCGCCACAGACGCTCAACATCTCGTGGAC 729
Qy 21 ThrGlySerSerAsnAspHeaVal 28
Db 730 ACGGCGAGTAGTAACTTGGAGTG 753
RESULT 12
AK046175 4101 bp mRNA linear HTC 03-Apr-2004
LOCUS Mus musculus adult male corpora quadrigemina cDNA, RIKEN
DEFINITION full-length enriched library, clone: B230346M13 product: beta-site
APC cleaving enzyme, full insert sequence.
ACCESSION AK046175 GI:26337868
VERSION AK046175.1
KEYWORDS HTC; CAP trapper.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS Carninci, P. and Hayashizaki, Y.
TITLE High-efficiency full-length cDNA cloning
JOURNAL Meth. Enzymol. 303, 19-44 (1999)
MEDLINE 99279253
PUBMED 10349636
AUTHORS 2
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Komano, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
TITLE Normalization and subtraction of cap-trapper-selected cDNAs to
JOURNAL prepare full-length cDNA libraries for rapid discovery of new genes
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AUTHORS 3
Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Komano, H., Akiyama, J., Nishi, K., Kitesuna, T., Tashiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K.,
Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M.,
Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsunura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE RIKEN integrated sequence analysis (RISA) system-384-format
JOURNAL RIKEN integrated sequence analysis (RISA) system-384-format
MEDLINE Genome Res. 10 (11), 1757-1771 (2000)
PUBMED 20530913
REFERENCES 11076861

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AUTHORS
4
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PANTOM Consortium.
TITLE
Functional annotation of a full-length mouse cDNA collection
JOURNAL
Nature 409, 685-690 (2001)
REFERENCE
AUTHORS
5
The PANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
JOURNAL
Nature 420, 563-573 (2002)
REFERENCE
AUTHORS
6 (bases 1 to 4101)
Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T.,
Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Kato, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M.,
Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M.,
Nakamura, M., Nishi, K., Nomura, K., Numata, K., Ohno, M., Ohnato, N.,
Okazaki, Y., Salto, R., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N.,
Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T.,
Sogabe, Y., Tagami, M., Tagawa, K., Takahashi, F., Takaku-Akahira, S.,
Takeda, Y., Tanaka, T., Tomaru, A., Toyota, T., Yasunishi, A.,
Muramatsu, M. and Hayashizaki, Y.
TITLE
Direct Submission
JOURNAL
Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
Physical and Chemical Research (RIKEN), Laboratory for Genome
Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
Kanagawa 230-0045, Japan (E-mail: genome-res@gs.c.riken.jp,
URL: http://genome.gsc.riken.jp/, Tel: 81-45-503-9222,
Fax: 81-45-503-9216)
COMMENT
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Please visit our web site for further details.
URL: http://genome.gsc.riken.jp/
URL: http://fantom.gsc.riken.jp/.
FEATURES
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Score: 144.00
Percent Similarity: 100.00%
Best Local Similarity: 100.00%
Query Match: 100.00%
DB: 3
Gaps: 0
US-10-726-967a-52 (1-28) x AK046175 (1-4101)
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666 GCGTACTATGTCGAGATATACCGTAGCGAGCCCCCAGACGCTCAGCATCTGCTGAC 725
Db 21 ThrGlySerSerAsnPhaIVal 28
726 ACGGCGAGTAGTAACCTTGACAGTG 749
RESULT 13
AK049626
LOCUS
DEFINITION
AK049626 4046 bp mRNA linear HTC 03-APR-2004
MUS musculus 12 days embryo spinal cord cDNA, RIKEN full-length
enriched library, clone: C530008K17 product: beta-site APP cleaving
enzyme, full insert sequence.
ACCESSION
AK049626
VERSION
AK049626.1 GI:26340361
KEYWORDS
HTC; CAP trapper.
SOURCE
MUS musculus (house mouse)
ORGANISM
Mus musculus
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS
Carninci, P. and Hayashizaki, Y.
TITLE
High-efficiency full-length cDNA cloning
JOURNAL
Meth. Enzymol. 303, 19-44 (1999)
MEDLINE
9279253
PUBMED
10349636
REFERENCE
AUTHORS
2
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
TITLE
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL
Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE
20499374
PUBMED
11042159
REFERENCE
AUTHORS
3
Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
Konno, H., Akiyama, J., Nishi, K., Kitsuana, T., Teshiro, H., Itoh, M.,
Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishino, T., Harada, A.,
Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kaishiwagi, K.,
Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M.,
Okazaki, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J.,
Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
TITLE
RIKEN integrated sequence analysis (RISA) system-384-format
sequencing pipeline with 384 multicapillary sequencer
JOURNAL
Genome Res. 10 (11), 1757-1771 (2000)
MEDLINE
20530913
PUBMED
11076861
REFERENCE
AUTHORS
4
The RIKEN Genome Exploration Research Group Phase II Team and the
PANTOM Consortium.
TITLE
Functional annotation of a full-length mouse cDNA collection
JOURNAL
Nature 409, 685-690 (2001)
REFERENCE
AUTHORS
5
The PANTOM Consortium and the RIKEN Genome Exploration Research
Group Phase I & II Team.
TITLE
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
JOURNAL
Nature 420, 563-573 (2002)
REFERENCE
AUTHORS
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Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
Hayashida, K., Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T.,
Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T.,
Kato, H., Kawai, J., Kojima, Y., Kondo, S., Konno, H., Kouda, M.,

ORIGIN
Alignment Scores:
Pred. No.:

3.4e-12 Length: 4101

Koya, S., Kuribara, C., Matsuyama, T., Miyazaki, A., Murata, M., Nakamura, M., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Ohnaka, N., Okazaki, Y., Saito, D., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, R., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Kahira, S., Takeeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A., Muramatsu, M., and Hayashizaki, Y.

TITLE
Direct Submission

JOURNAL
Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp, URL: http://genome.gsc.riken.jp/, Tel: 01-45-503-9222, Fax: 01-45-503-9216)

COMMENT
cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. Please visit our web site for further details.
URL: http://genome.gsc.riken.jp/
URL: http://fantom.gsc.riken.jp/
Location/Qualifiers

FEATURES
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ORIGIN

Alignment Scores:

Pred. No.:	4.9e-12	Length: 27
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Percent Similarity:	100.00%	Conservative: 1
Best Local Similarity:	96.43%	Mismatches: 0
Query Match:	99.31%	Indels: 0
DB:	3	Gaps: 0

US-10-726-967a-52 (1-28) x AK049626 (1-4046)

Qy 1 G1YTYTYValGluMetThrValGlySerProFrogInThrLeuAniLeuValasp 20
|||||
Db 652 GGGTACTAGTGGAGATGACCATGCGAGCCCCCAGACGCTCAACATCTGTGTGAC 711
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Qy 21 ThG1YserSerAnphea1aVal 28
|||||
Db 712 ACCGGCAGTAGTAACTTGGCAGTG 735
|||||

RESULT 14
AL700831

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
MIPS
Ingolstaedter Landstr.1, D-85764 Neuberg, Germany
This is the 5' sequence of the clone insert
clone from S. Wiemann, Molecular Genome Analysis, German Cancer
Research Center (DKFZ), Email: s.wiemann@dkfz-heidelberg.de;
sequenced by GBF (National Research Centre for Biotechnology Ltd.,
Braunschweig/Germany) within the cDNA sequencing consortium of the
German Genome Project.
No 5' sequence available.
This clone (DKFZ68612411) is available at the RZPD in Berlin.
Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059
Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.

FEATURES
source
1. 461
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cDNA-collection"

ORIGIN

Alignment Scores:

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Percent Similarity:	96.43%	Conservative: 0
Best Local Similarity:	96.43%	Mismatches: 1
Query Match:	95.83%	Indels: 0
DB:	1	Gaps: 0

US-10-726-967a-52 (1-28) x AL700831 (1-461)

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Qy 21 ThG1YserSerAnphea1aVal 28
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Db 130 ACCGGCAGTAGTAACTTGGCAGTG 153
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RESULT 15
BG288435
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
1 (bases 1 to 720)

AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
 TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabbs-remail.nih.gov

Tissue Procurement: ATCC
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
 DNA Sequencing by: Incyte Genomics, Inc.
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LNL at:
<http://image.llnl.gov>
 Plate: LLM10367 row: h column: 02
 High quality sequence start: 8
 High quality sequence stop: 715.
 Location/Qualifiers
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 /organism="Mus musculus"
 /mol_type="mRNA"
 /db_xref="taxon:10090"
 /clone="IMAGE:4500913"
 /tissue_type="retina"
 /lab_host="DH10B (phage-resistant)"
 /clone_id="NIH_MGC_94"
 /note="Organ: eye; Vector: pCMV-SPORT6; Site_1: NotI;
 Site_2: SalI; Cloned unidirectionally; oligo-dT primed.
 Average insert size 3.3 kb. Library enriched for
 full-length clones and constructed by Life Technologies.
 Note: this is a NIH_MGC library."

FEATURES

source

ORIGIN

Alignment Scores:

Pred. No.:	4.05e-12	Length:	720
Score:	138.00	Matches:	27
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	95.83%	Indels:	0
DB:	4	Gaps:	0

US-10-726-967a-52 (1-28) x BG288435 (1-720)

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DB	8	TACTATGTGAGATGACCGTAGGCAAGCCGCCACAGACCGCTCAACATCTGTGTGACACG	67
QY	22	GlySerSerAsnPheAlaVal	28
DB	68	GGCAGTAGTAACTTGCAAGTG	88

Search completed: July 27, 2005, 18:17:56
 Job time : 3160 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: July 26, 2005, 16:38:55 ; Search time 41 Seconds
(without alignments)
50.980 Million cell updates/sec

Title: US-10-726-967A-52

Perfect score: 144

Sequence: 1 GYVEMTVGSPQTLNLTVDGSSNPAV 28

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 513545 seqs, 74649064 residues

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents AA:*

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3: /cgnt2_6/prodata/1/iaa/5A_COMB.pep:.*
4: /cgnt2_6/prodata/1/iaa/5B_COMB.pep:.*
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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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2	144	100.0	361	4 US-09-724-566A-75	Sequence 75, App1
3	144	100.0	361	4 US-09-471-669A-75	Sequence 75, App1
4	144	100.0	374	4 US-09-724-566A-71	Sequence 71, App1
5	144	100.0	374	4 US-09-471-669A-71	Sequence 71, App1
6	144	100.0	360	4 US-09-471-669A-108	Sequence 108, App
7	144	100.0	390	4 US-09-724-566A-70	Sequence 70, App1
8	144	100.0	390	4 US-09-471-669A-70	Sequence 70, App1
9	144	100.0	395	4 US-09-724-566A-68	Sequence 68, App1
10	144	100.0	395	4 US-09-471-669A-68	Sequence 68, App1
11	144	100.0	401	4 US-09-471-669A-106	Sequence 106, App
12	144	100.0	407	4 US-09-724-566A-58	Sequence 58, App1
13	144	100.0	407	4 US-09-471-669A-58	Sequence 58, App1
14	144	100.0	408	4 US-09-471-669A-105	Sequence 105, App
15	144	100.0	419	4 US-09-724-566A-57	Sequence 57, App1
16	144	100.0	419	4 US-09-471-669A-57	Sequence 57, App1
17	144	100.0	420	4 US-09-724-566A-60	Sequence 60, App1
18	144	100.0	420	4 US-09-471-669A-60	Sequence 60, App1
19	144	100.0	425	4 US-09-548-372D-28	Sequence 28, App1
20	144	100.0	425	4 US-09-548-372D-28	Sequence 28, App1
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24	144	100.0	425	4 US-09-794-927A-28	Sequence 28, App1
25	144	100.0	425	4 US-09-548-373D-28	Sequence 28, App1
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27	144	100.0	425	4 US-09-869-414-28	Sequence 28, App1

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35	144	100.0	428	4 US-09-551-853D-51	Sequence 51, App1
36	144	100.0	428	4 US-09-416-901B-51	Sequence 51, App1
37	144	100.0	428	4 US-09-548-376D-51	Sequence 51, App1
38	144	100.0	428	4 US-09-794-927A-51	Sequence 51, App1
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ALIGNMENTS

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RESULT 1
US-09-471-669A-107
; Sequence 107, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basl, Guridgal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Srinna, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McCollough, Lisa
; APPLICANT: Eian Pharmaceuticals, Inc.
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
; FILE REFERENCE: 015270-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; CURRENT FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 107
; LENGTH: 231
; TYPE: PRT
; ORGANISM: Mus sp.
; OTHER INFORMATION: PBS/MultiPain E17 Brain #17 construct
US-09-471-669A-107

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Best Local Similarity 100.0%; Pred. No. 6.5e-14;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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RESULT 2
US-09-724-566A-75
; Sequence 75, Application US/09724566A
; Patent No. 6627739
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; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6627739mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
; FILE REFERENCE: 228-US-NEWC2
; CURRENT APPLICATION NUMBER: US/09/724,566A
; PRIOR FILING DATE: 2000-11-28
; PRIOR APPLICATION NUMBER: US 09/501,708
; PRIOR FILING DATE: 2000-02-10
; PRIOR APPLICATION NUMBER: 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: 60/139,172
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 75
; LENGTH: 361
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-724-566A-75

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Best Local Similarity 100.0%; Score 144; DB 4; Length 361;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 12 GYVEMTVGSPPTNLTIVDTGSSNFAV 39

RESULT 3
US-09-471-669A-75
; Sequence 75, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; APPLICANT: Elan Pharmaceuticals, Inc.
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
; FILE REFERENCE: 015770-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; PRIOR FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 75
; LENGTH: 361
; TYPE: PRT
; ORGANISM: Homo sapiens
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US-09-471-669A-75

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Best Local Similarity 100.0%; Score 144; DB 4; Length 361;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 12 GYVEMTVGSPPTNLTIVDTGSSNFAV 39

RESULT 4
US-09-724-566A-71
; Sequence 71, Application US/09724566A
; Patent No. 6627739
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6627739mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
; FILE REFERENCE: 228-US-NEWC2
; CURRENT APPLICATION NUMBER: US/09/724,566A
; PRIOR FILING DATE: 2000-11-28
; PRIOR APPLICATION NUMBER: US 09/501,708
; PRIOR FILING DATE: 2000-02-10
; PRIOR APPLICATION NUMBER: 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: 60/139,172
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 71
; LENGTH: 374
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-724-566A-71

Query Match
Best Local Similarity 100.0%; Score 144; DB 4; Length 374;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GYVEMTVGSPPTNLTIVDTGSSNFAV 28
Db 29 GYVEMTVGSPPTNLTIVDTGSSNFAV 56

RESULT 5
US-09-471-669A-71
; Sequence 71, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; APPLICANT: Elan Pharmaceuticals, Inc.
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
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FILE REFERENCE: 015270-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; CURRENT FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 71
; LENGTH: 374
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-471-669A-71

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Best Local Similarity 100.0%; Pred. No. 1,1e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 29 GYVENTVGSPPQTINILVDTGSSNPAV 56

RESULT 6
US-09-471-669A-108
; Sequence 108, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; APPLICANT: Elan Pharmaceuticals, Inc.
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
; FILE REFERENCE: 015270-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; CURRENT FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 108
; LENGTH: 380
; TYPE: PRT
; ORGANISM: Mus sp.
; FEATURE:
; OTHER INFORMATION: pbs/Multipain E17 Brain#15 construct
; US-09-471-669A-108

Query Match          100.0%; Score 144; DB 4; Length 380;
Best Local Similarity 100.0%; Pred. No. 1,2e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GYVENTVGSPPQTINILVDTGSSNPAV 28
Db 15 GYVENTVGSPPQTINILVDTGSSNPAV 42

RESULT 7
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US-09-724-566A-70
; Sequence 70, Application US/09724566A
; Patent No. 6627739
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6627739mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
; FILE REFERENCE: 228-US-NEWC2
; CURRENT APPLICATION NUMBER: US/09/724,566A
; CURRENT FILING DATE: 2000-11-28
; PRIOR APPLICATION NUMBER: US 09/501,708
; PRIOR FILING DATE: 2000-02-10
; PRIOR APPLICATION NUMBER: 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 70
; LENGTH: 390
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-724-566A-70

Query Match          100.0%; Score 144; DB 4; Length 390;
Best Local Similarity 100.0%; Pred. No. 1,2e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GYVENTVGSPPQTINILVDTGSSNPAV 28
Db 12 GYVENTVGSPPQTINILVDTGSSNPAV 39

RESULT 8
US-09-471-669A-70
; Sequence 70, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; APPLICANT: Elan Pharmaceuticals, Inc.
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
; FILE REFERENCE: 015270-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; CURRENT FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 70
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LENGTH: 390
TYPE: PRT
ORGANISM: Homo sapiens
US-09-471-669A-70

Query Match
Best Local Similarity 100.0%; Score 144; DB 4; Length 390;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLITLVDGSSNFAY 28
DB 12 GYVEMTVGSPPTLITLVDGSSNFAY 39

RESULT 9
US-09-724-566A-68

Sequence 68, Application US/09724566A
Patent No. 6627739

GENERAL INFORMATION:

APPLICANT: Anderson, John P.

APPLICANT: Baal, Gurigbal

APPLICANT: Doane, Minh Tam

APPLICANT: Frigon, No. 6627739mand

APPLICANT: John, Varghese

APPLICANT: Power, Michael

APPLICANT: Sinha, Sukanto

APPLICANT: Tatsuno, Gwen

APPLICANT: Tung, Jay

APPLICANT: Wang, Shuwen

APPLICANT: McConlogue, Lisa

TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and

FILE REFERENCE: 228-US-NEMC2

CURRENT APPLICATION NUMBER: US/09/724,566A

PRIOR FILING DATE: 2000-11-28

PRIOR APPLICATION NUMBER: US 09/501,708

PRIOR FILING DATE: 2000-02-10

PRIOR APPLICATION NUMBER: 60/119,571

PRIOR FILING DATE: 1999-02-10

PRIOR APPLICATION NUMBER: 60/139,172

PRIOR FILING DATE: 1999-06-15

NUMBER OF SEQ ID NOS: 104

SOFTWARE: PaateSeq for Windows Version 4.0

SEQ ID NO 68

LENGTH: 395

TYPE: PRT

ORGANISM: Homo sapiens

US-09-724-566A-68

Query Match

Best Local Similarity 100.0%; Score 144; DB 4; Length 395;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLITLVDGSSNFAY 28
DB 17 GYVEMTVGSPPTLITLVDGSSNFAY 44

RESULT 10
US-09-471-669A-68

Sequence 68, Application US/09471669A
Patent No. 6830918

GENERAL INFORMATION:

APPLICANT: Anderson, John P.

APPLICANT: Baal, Gurigbal

APPLICANT: Doane, Minh Tam

APPLICANT: Frigon, No. 6830918mand

APPLICANT: John, Varghese

APPLICANT: Power, Michael

APPLICANT: Sinha, Sukanto

APPLICANT: Tatsuno, Gwen

APPLICANT: Tung, Jay

APPLICANT: Wang, Shuwen

APPLICANT: McConlogue, Lisa
APPLICANT: Brian Pharmaceuticals, Inc.
TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
FILE REFERENCE: 015270-006430US

CURRENT APPLICATION NUMBER: US/09/471,669A

PRIOR FILING DATE: 1999-12-24

PRIOR APPLICATION NUMBER: US 60/114,408

PRIOR FILING DATE: 1998-12-31

PRIOR APPLICATION NUMBER: US 60/119,571

PRIOR FILING DATE: 1999-02-10

PRIOR APPLICATION NUMBER: US 60/139,172

PRIOR FILING DATE: 1999-06-15

NUMBER OF SEQ ID NOS: 108

SOFTWARE: PatentIn Ver. 2.1

SEQ ID NO 68

LENGTH: 395

TYPE: PRT

ORGANISM: Homo sapiens

US-09-471-669A-68

Query Match

Best Local Similarity 100.0%; Score 144; DB 4; Length 395;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLITLVDGSSNFAY 28
DB 17 GYVEMTVGSPPTLITLVDGSSNFAY 44

RESULT 11
US-09-471-669A-106

Sequence 106, Application US/09471669A
Patent No. 6830918

GENERAL INFORMATION:

APPLICANT: Anderson, John P.

APPLICANT: Baal, Gurigbal

APPLICANT: Doane, Minh Tam

APPLICANT: Frigon, No. 6830918mand

APPLICANT: John, Varghese

APPLICANT: Power, Michael

APPLICANT: Sinha, Sukanto

APPLICANT: Tatsuno, Gwen

APPLICANT: Tung, Jay

APPLICANT: Wang, Shuwen

APPLICANT: McConlogue, Lisa

TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS

FILE REFERENCE: 015270-006430US

CURRENT APPLICATION NUMBER: US/09/471,669A

PRIOR FILING DATE: 1999-12-24

PRIOR APPLICATION NUMBER: US 60/114,408

PRIOR FILING DATE: 1998-12-31

PRIOR APPLICATION NUMBER: US 60/119,571

PRIOR FILING DATE: 1999-02-10

PRIOR APPLICATION NUMBER: US 60/139,172

PRIOR FILING DATE: 1999-06-15

NUMBER OF SEQ ID NOS: 108

SOFTWARE: PatentIn Ver. 2.1

SEQ ID NO 106

LENGTH: 401

TYPE: PRT

ORGANISM: Mus sp.

OTHER INFORMATION: pbs/MuImPain E17 #14 construct

US-09-471-669A-106

Query Match

Best Local Similarity 100.0%; Score 144; DB 4; Length 401;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLITLVDGSSNFAY 28
DB 25 GYVEMTVGSPPTLITLVDGSSNFAY 52


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RESULT 12
US-09-724-566A-58
; Sequence 58, Application US/09724566A
; Patent No. 6627739
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6627739mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
; TITLE OF INVENTION: Methods
; FILE REFERENCE: 228-US-NEWC2
; CURRENT APPLICATION NUMBER: US/09/724,566A
; PRIOR FILING DATE: 2000-11-28
; PRIOR APPLICATION NUMBER: US 09/501,708
; PRIOR FILING DATE: 2000-02-10
; PRIOR APPLICATION NUMBER: 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 58
; LENGTH: 407
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-724-566A-58

Query Match          100.0%; Score 144; DB 4; Length 407;
Best Local Similarity 100.0%; Pred. No. 1.3e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GYVETVGSPPQTNIILVDTGSSNPAV 28
DB      29 GYVETVGSPPQTNIILVDTGSSNPAV 56

RESULT 13
US-09-471-669A-58
; Sequence 58, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
; FILE REFERENCE: 015270-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; PRIOR FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 105
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 105
; LENGTH: 408
; TYPE: PRT
; ORGANISM: Mus sp.
; FEATURE:
; OTHER INFORMATION: PBS/Multipain E17 #11 construct
US-09-471-669A-105

Query Match          100.0%; Score 144; DB 4; Length 408;
Best Local Similarity 100.0%; Pred. No. 1.3e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GYVETVGSPPQTNIILVDTGSSNPAV 28
DB      27 GYVETVGSPPQTNIILVDTGSSNPAV 54

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; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 58
; LENGTH: 407
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-471-669A-58

Query Match          100.0%; Score 144; DB 4; Length 407;
Best Local Similarity 100.0%; Pred. No. 1.3e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GYVETVGSPPQTNIILVDTGSSNPAV 28
DB      29 GYVETVGSPPQTNIILVDTGSSNPAV 56

RESULT 14
US-09-471-669A-105
; Sequence 105, Application US/09471669A
; Patent No. 6830918
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6830918mand
; APPLICANT: John, Varghese
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatsuno, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; APPLICANT: Eilan Pharmaceuticals, Inc.
; TITLE OF INVENTION: BETA-SECRETASE ENZYME COMPOSITIONS AND METHODS
; FILE REFERENCE: 015270-006430US
; CURRENT APPLICATION NUMBER: US/09/471,669A
; PRIOR FILING DATE: 1999-12-24
; PRIOR APPLICATION NUMBER: US 60/114,408
; PRIOR FILING DATE: 1998-12-31
; PRIOR APPLICATION NUMBER: US 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: US 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 108
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 105
; LENGTH: 408
; TYPE: PRT
; ORGANISM: Mus sp.
; FEATURE:
; OTHER INFORMATION: PBS/Multipain E17 #11 construct
US-09-471-669A-105

Query Match          100.0%; Score 144; DB 4; Length 408;
Best Local Similarity 100.0%; Pred. No. 1.3e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GYVETVGSPPQTNIILVDTGSSNPAV 28
DB      27 GYVETVGSPPQTNIILVDTGSSNPAV 54

RESULT 15
US-09-724-566A-57
; Sequence 57, Application US/09724566A
; Patent No. 6627739
; GENERAL INFORMATION:
; APPLICANT: Anderson, John P.
; APPLICANT: Basi, Gurigbal
; APPLICANT: Doane, Minh Tam
; APPLICANT: Frigon, No. 6627739mand
; APPLICANT: John, Varghese

```

```
; APPLICANT: Power, Michael
; APPLICANT: Sinha, Sukanto
; APPLICANT: Tatarsko, Gwen
; APPLICANT: Tung, Jay
; APPLICANT: Wang, Shuwen
; APPLICANT: McConlogue, Lisa
; TITLE OF INVENTION: Beta-Secretase Enzyme Compositions and
; TITLE OF INVENTION: Methods
; FILE REFERENCE: 228-US-NEMC2
; CURRENT APPLICATION NUMBER: US/09/724,566A
; CURRENT FILING DATE: 2000-11-28
; PRIOR APPLICATION NUMBER: US 09/501,708
; PRIOR FILING DATE: 2000-02-10
; PRIOR APPLICATION NUMBER: 60/119,571
; PRIOR FILING DATE: 1999-02-10
; PRIOR APPLICATION NUMBER: 60/139,172
; PRIOR FILING DATE: 1999-06-15
; NUMBER OF SEQ ID NOS: 104
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 57
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-724-566A-57
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Query Match 100.0%; Score 144; DB 4; Length 419;
Best Local Similarity 100.0%; Pred. No. 1.3e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
   |||||
Db 74 GYVEMTVGSPPTLNIIVDTGSSNFAV 101
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Search completed: July 26, 2005, 16:48:45
Job time : 41 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd

OM protein - protein search, using sw model

Run on: July 26, 2005, 16:41:36 ; Search time 154 Seconds
(without alignments)
70.726 Million cell updates/sec

Title:	US-10-726-967A-52
Perfect score:	144
Sequence:	I GYVEMTVGSPQTLNILVDTGSSNFAV 28

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

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Minimum DB seq length: 0
Maximum DB seq length: 2000000000
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Post-processing:	Minimum Match	0%
	Maximum Match	100%
	Listing first	45 summaries

Published applications: AA: *

- 1: /cgn2_6/ptodata/2/pubpaa/US07_PUBCOMB.pdp: *
- 2: /cgn2_6/ptodata/2/pubpaa/PCT_NEW_PUB.pdp: *
- 3: /cgn2_6/ptodata/2/pubpaa/US06_NEW_PUB.pdp: *
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- 8: /cgn2_6/ptodata/2/pubpaa/US09_PUBCOMB.pdp: *
- 9: /cgn2_6/ptodata/2/pubpaa/US09A_PUBCOMB.pdp: *
- 10: /cgn2_6/ptodata/2/pubpaa/US09C_PUBCOMB.pdp: *
- 11: /cgn2_6/ptodata/2/pubpaa/US09C_PUBCOMB.pdp: *
- 12: /cgn2_6/ptodata/2/pubpaa/US09_NEW_PUB.pdp: *
- 13: /cgn2_6/ptodata/2/pubpaa/US10A_PUBCOMB.pdp: *
- 14: /cgn2_6/ptodata/2/pubpaa/US10B_PUBCOMB.pdp: *
- 15: /cgn2_6/ptodata/2/pubpaa/US10C_PUBCOMB.pdp: *
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- 18: /cgn2_6/ptodata/2/pubpaa/US11A_PUBCOMB.pdp: *
- 19: /cgn2_6/ptodata/2/pubpaa/US11A_PUBCOMB.pdp: *
- 20: /cgn2_6/ptodata/2/pubpaa/US11_NEW_PUB.pdp: *
- 21: /cgn2_6/ptodata/2/pubpaa/US60_NEW_PUB.pdp: *
- 22: /cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pdp: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result No	Score	Query Match	length	DB	ID	Description
1	144	100.0	28	17	US-10-726-9678-52	Sequence 52, App
2	144	100.0	358	17	US-10-872-198-12	Sequence 12, App
3	144	100.0	358	17	US-10-872-1978-12	Sequence 12, App
4	144	100.0	331	15	US-10-372-473-4	Sequence 4, App1
5	144	100.0	403	15	US-10-400-273-4	Sequence 4, App1
6	144	100.0	406	16	US-10-837-021A-2	Sequence 2, App1
7	144	100.0	406	16	US-10-837-021A-3	Sequence 3, App1
8	144	100.0	406	16	US-10-837-021A-4	Sequence 4, App1
9	144	100.0	406	16	US-10-837-021A-5	Sequence 5, App1
10	144	100.0	408	15	US-10-400-273-5	Sequence 5, App1
11	144	100.0	411	15	US-10-400-273-1	Sequence 1, App1

ALIGNMENTS

12	144	100.0	411	15	US-10-627-473-19	Sequence 19, App
13	144	100.0	411	16	US-10-627-473-20	Sequence 20, App
14	144	100.0	414	15	US-10-281-092-9	Sequence 9, App
15	144	100.0	417	15	US-10-627-473-21	Sequence 21, App
16	144	100.0	425	9	US-09-794-927-28	Sequence 28, App
17	144	100.0	425	9	US-09-795-447-28	Sequence 28, App
18	144	100.0	425	9	US-09-794-743-28	Sequence 28, App
19	144	100.0	425	9	US-09-794-428-28	Sequence 28, App
20	144	100.0	425	9	US-09-794-925-28	Sequence 28, App
21	144	100.0	425	9	US-09-681-442-28	Sequence 28, App
22	144	100.0	425	10	US-09-869-414-28	Sequence 28, App
23	144	100.0	425	10	US-09-548-366-28	Sequence 28, App
24	144	100.0	425	15	US-10-372-473-3	Sequence 3, App
25	144	100.0	425	15	US-10-652-927-28	Sequence 28, App
26	144	100.0	425	15	US-10-652-830-28	Sequence 28, App
27	144	100.0	425	16	US-10-652-935-28	Sequence 28, App
28	144	100.0	425	16	US-10-476-935-28	Sequence 28, App
29	144	100.0	425	17	US-10-940-867-28	Sequence 28, App
30	144	100.0	425	17	US-10-477-076-28	Sequence 28, App
31	144	100.0	428	9	US-09-794-927-51	Sequence 51, App
32	144	100.0	428	9	US-09-795-847-51	Sequence 51, App
33	144	100.0	428	9	US-09-794-743-51	Sequence 51, App
34	144	100.0	428	9	US-09-794-748-51	Sequence 51, App
35	144	100.0	428	9	US-09-794-925-51	Sequence 51, App
36	144	100.0	428	9	US-09-681-442-51	Sequence 51, App
37	144	100.0	428	10	US-09-869-414-51	Sequence 51, App
38	144	100.0	428	10	US-09-548-366-51	Sequence 51, App
39	144	100.0	428	15	US-10-652-927-51	Sequence 51, App
40	144	100.0	428	15	US-10-652-830-51	Sequence 51, App
41	144	100.0	428	16	US-10-652-945-51	Sequence 51, App
42	144	100.0	428	16	US-10-476-935-51	Sequence 51, App
43	144	100.0	428	17	US-10-477-076-51	Sequence 51, App
44	144	100.0	432	15	US-10-372-473-2	Sequence 26, App
45	144	100.0	433	9	US-09-794-927-26	Sequence 26, App

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RESULT 1
US-10-726-967A-52
; Sequence 52, Application US/10726967A
; Publication No. US20050074456A1
GENERAL INFORMATION:
APPLICANT: Ballinger, Marcus
TITLE OF INVENTION: Constructs for Homogenously Processed Preparations of Beta Site
TITLE OF INVENTION: App-Cleaving Enzyme
FILE REFERENCE: 2004345-0021
CURRENT APPLICATION NUMBER: US/10/726,967A
CURRENT FILING DATE: 2003-12-02
NUMBER OF SEQ ID NOS: 110
SOFTWARE: PatentIn version 3.2
SEQ ID NO 52
LENGTH: 28
TYPE: PRT
ORGANISM: Artificial
FEATURES:
OTHER INFORMATION: Residues 74-101 of human BACE1 preprosequence
US-10-726-967A-52

Query Match          100.0%; Score 144; DB 17; Length 28;
Best Local Similarity 100.0%; Pred.No. 1.8e-14;
Matches      28; Conservative    0; Mismatches     0; Gaps      0; Indels      0;

Oy              1 GYVEMTVGSPPTTLNIIIVDTGSSNFAY 28
                |||||
Db              1 GYVEMTVGSPPTTLNIIIVDTGSSNFAY 28

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/ GENERAL INFORMATION:
/ APPLICANT: Ulrich HAUPTS
/ APPLICANT: Andre KOLTERMANN
/ APPLICANT: Andreas SCHEIDIG
/ APPLICANT: Christian VOTSMERER
/ APPLICANT: Ulrich KETTLING
/ TITLE OF INVENTION: NEW BIOLOGICAL ENTITIES AND USE THEREOF
/ FILE REFERENCE: 04156.0002U4
/ CURRENT APPLICATION NUMBER: US/10/872,198
/ CURRENT FILING DATE: 2004-06-18
/ PRIOR APPLICATION NUMBER: 60/543,518
/ PRIOR FILING DATE: 2004-02-11
/ PRIOR APPLICATION NUMBER: 60/524,960
/ PRIOR FILING DATE: 2003-11-25
/ PRIOR APPLICATION NUMBER: EP 04003058
/ PRIOR FILING DATE: 2004-02-11
/ PRIOR APPLICATION NUMBER: EP 03025871
/ PRIOR FILING DATE: 2003-11-11
/ PRIOR APPLICATION NUMBER: EP 03025851
/ PRIOR FILING DATE: 2003-11-10
/ PRIOR APPLICATION NUMBER: EP 03013819
/ PRIOR FILING DATE: 2003-06-18
/ NUMBER OF SEQ ID NOS: 149
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 12
/ LENGTH: 358
/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-10-872-198-12
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Query Match      100.0%; Score 144; DB 17; Length 358;
Best Local Similarity 100.0%; Pred. No. 3,5e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      1 GYVEMTVGSPPTNTILVDTGSSNFAV 28
      |||
      13 GYVEMTVGSPPTNTILVDTGSSNFAV 40
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RESULT 3
US-10-872-197A-12
/ Sequence 12, Application US/10872197A
/ Publication No. US20050059126A1
/ GENERAL INFORMATION:
/ APPLICANT: Ulrich HAUPTS
/ APPLICANT: Andre KOLTERMANN
/ APPLICANT: Andreas SCHEIDIG
/ APPLICANT: Christian VOTSMERER
/ APPLICANT: Ulrich KETTLING
/ TITLE OF INVENTION: NEW BIOLOGICAL ENTITIES AND USE THEREOF
/ FILE REFERENCE: 04156.0002U3
/ CURRENT APPLICATION NUMBER: US/10/872,197A
/ CURRENT FILING DATE: 2004-06-18
/ PRIOR APPLICATION NUMBER: 60/524,960
/ PRIOR FILING DATE: 2003-11-25
/ PRIOR APPLICATION NUMBER: EP 03025871
/ PRIOR FILING DATE: 2003-11-11
/ PRIOR APPLICATION NUMBER: EP 03013819
/ PRIOR FILING DATE: 2003-11-10
/ NUMBER OF SEQ ID NOS: 96
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 12
/ LENGTH: 358
/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-10-872-197A-12
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Query Match      100.0%; Score 144; DB 17; Length 358;
Best Local Similarity 100.0%; Pred. No. 3,5e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db      13 GYVEMTVGSPPTNTILVDTGSSNFAV 40
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RESULT 4
US-10-372-473-4
/ Sequence 4, Application US/10372473
/ Publication No. US20040005691A1
/ GENERAL INFORMATION:
/ APPLICANT: Chou, Kuo-Chen
/ APPLICANT: Howe, W. Jeffery
/ TITLE OF INVENTION: Modified BACE
/ FILE REFERENCE: MBHB-01-1766-A
/ CURRENT APPLICATION NUMBER: US/10/372,473
/ CURRENT FILING DATE: 2003-02-21
/ NUMBER OF SEQ ID NOS: 24
/ SOFTWARE: PatentIn version 3.2
/ SEQ ID NO 4
/ LENGTH: 391
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ FEATURE:
/ NAME/KEY: MISC FEATURE
/ OTHER INFORMATION: Human beta-secretase.
US-10-372-473-4
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Query Match      100.0%; Score 144; DB 15; Length 391;
Best Local Similarity 100.0%; Pred. No. 3,9e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      1 GYVEMTVGSPPTNTILVDTGSSNFAV 28
      |||
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RESULT 5
US-10-400-273-4
/ Sequence 4, Application US/10400273
/ Publication No. US20040014194A1
/ GENERAL INFORMATION:
/ APPLICANT: Beyer, Brian
/ APPLICANT: Hammond, Gerald S
/ APPLICANT: Reichert, Paul
/ APPLICANT: Strickland, Corey
/ APPLICANT: Wang, Wenyan
/ APPLICANT: Weber, Patricia C
/ APPLICANT: Wong, Gwendolyn
/ APPLICANT: Zhang, Lili
/ TITLE OF INVENTION: BETA-SECRETASE CRYSTALS AND METHODS FOR PREPARING AND USING THE S.
/ FILE REFERENCE: JBO1531-K-US
/ CURRENT APPLICATION NUMBER: US/10/400,273
/ CURRENT FILING DATE: 2003-03-26
/ PRIOR APPLICATION NUMBER: 60/367,937
/ PRIOR FILING DATE: 2002-03-27
/ NUMBER OF SEQ ID NOS: 5
/ SOFTWARE: PatentIn version 3.1
/ SEQ ID NO 4
/ LENGTH: 403
/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-10-400-273-4
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Query Match      100.0%; Score 144; DB 15; Length 403;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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```
QY      1 GYVEMTVGSPPTNTILVDTGSSNFAV 28
      |||
      27 GYVEMTVGSPPTNTILVDTGSSNFAV 54
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RESULT 6
US-10-837-021A-2
/ Sequence 2, Application US/10837021A
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Publication No. US20040265965A1
GENERAL INFORMATION:
APPLICANT: Anderson, John P
APPLICANT: McConlogue, Lisa
APPLICANT: Basl, Guribdal
APPLICANT: Sinha, Sukarno
TITLE OF INVENTION: Glycosylation Variants of BACE
FILE REFERENCE: MBHB-03-268-A
CURRENT APPLICATION NUMBER: US/10/837,021A
PRIOR FILING DATE: 2004-04-30
PRIOR APPLICATION NUMBER: 60/467,509
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.3
SEQ ID NO 2
LENGTH: 406
TYPE: PRT
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Human BACE with asparagine to alanine (N223A) mutation.
US-10-837-021A-2

Query Match          100.0%; Score 144; DB 16; Length 406;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLNILVDTGSSNFAV 28
DB 29 GYVEMTVGSPPTLNILVDTGSSNFAV 56

RESULT 7
US-10-837-021A-3
Sequence 3, Application US/10837021A
Publication No. US20040265965A1
GENERAL INFORMATION:
APPLICANT: Anderson, John P
APPLICANT: McConlogue, Lisa
APPLICANT: Basl, Guribdal
APPLICANT: Sinha, Sukarno
TITLE OF INVENTION: Glycosylation Variants of BACE
FILE REFERENCE: MBHB-03-268-A
CURRENT APPLICATION NUMBER: US/10/837,021A
PRIOR FILING DATE: 2004-04-30
PRIOR APPLICATION NUMBER: 60/467,509
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.3
SEQ ID NO 3
LENGTH: 406
TYPE: PRT
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Human BACE with serine to isoleucine (S174I) and asparagine to
US-10-837-021A-3

Query Match          100.0%; Score 144; DB 16; Length 406;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLNILVDTGSSNFAV 28
DB 29 GYVEMTVGSPPTLNILVDTGSSNFAV 56

RESULT 8
US-10-837-021A-4
Sequence 4, Application US/10837021A
Publication No. US20040265965A1
GENERAL INFORMATION:
APPLICANT: Anderson, John P
APPLICANT: McConlogue, Lisa
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APPLICANT: Basl, Guribdal
APPLICANT: Sinha, Sukarno
TITLE OF INVENTION: Glycosylation Variants of BACE
FILE REFERENCE: MBHB-03-268-A
CURRENT APPLICATION NUMBER: US/10/837,021A
PRIOR FILING DATE: 2004-04-30
PRIOR APPLICATION NUMBER: 60/467,509
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.3
SEQ ID NO 4
LENGTH: 406
TYPE: PRT
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Human BACE with serine to isoleucine (S174I), asparagine to
US-10-837-021A-4

Query Match          100.0%; Score 144; DB 16; Length 406;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLNILVDTGSSNFAV 28
DB 29 GYVEMTVGSPPTLNILVDTGSSNFAV 56

RESULT 9
US-10-837-021A-5
Sequence 5, Application US/10837021A
Publication No. US20040265965A1
GENERAL INFORMATION:
APPLICANT: Anderson, John P
APPLICANT: McConlogue, Lisa
APPLICANT: Basl, Guribdal
APPLICANT: Sinha, Sukarno
TITLE OF INVENTION: Glycosylation Variants of BACE
FILE REFERENCE: MBHB-03-268-A
CURRENT APPLICATION NUMBER: US/10/837,021A
PRIOR FILING DATE: 2004-04-30
PRIOR APPLICATION NUMBER: 60/467,509
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.3
SEQ ID NO 5
LENGTH: 406
TYPE: PRT
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Human BACE with serine to isoleucine (S174I), asparagine to
US-10-837-021A-5

Query Match          100.0%; Score 144; DB 16; Length 406;
Best Local Similarity 100.0%; Pred. No. 4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLNILVDTGSSNFAV 28
DB 29 GYVEMTVGSPPTLNILVDTGSSNFAV 56

RESULT 10
US-10-400-273-5
Sequence 5, Application US/10400273
Publication No. US2004040014194A1
GENERAL INFORMATION:
APPLICANT: Bayer, Brian
APPLICANT: Hammond, Gerald S
APPLICANT: Reichert, Paul
APPLICANT: Strickland, Corey
```

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; APPLICANT: Wang, Wenyan
; APPLICANT: Weber, Patricia C
; APPLICANT: Wong, Gwendolyn
; APPLICANT: Zhang, Lili
; TITLE OF INVENTION: BETA-SECRETASE CRYSTALS AND METHODS FOR PREPARING AND USING THE S
; FILE REFERENCE: JB01531-K-US
; CURRENT APPLICATION NUMBER: US/10/400,273
; CURRENT FILING DATE: 2003-03-26
; PRIOR APPLICATION NUMBER: 60/367,937
; PRIOR FILING DATE: 2002-03-27
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 408
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-400-273-5

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Query Match          100.0%; Score 144; DB 15; Length 408;
Best Local Similarity 100.0%; Pred. No. 4,1e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
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Db 32 GYVEMTVGSPPTLNIIVDTGSSNFAV 59

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RESULT 11
US-10-400-273-1
; Sequence 1, Application US/10400273
; Publication No. US2004001494A1
; GENERAL INFORMATION:
; APPLICANT: Beyer, Brian
; APPLICANT: Hammond, Gerald S
; APPLICANT: Reichert, Paul
; APPLICANT: Strickland, Corey
; APPLICANT: Wang, Wenyan
; APPLICANT: Weber, Patricia C
; APPLICANT: Wong, Gwendolyn
; APPLICANT: Zhang, Lili
; TITLE OF INVENTION: BETA-SECRETASE CRYSTALS AND METHODS FOR PREPARING AND USING THE S
; FILE REFERENCE: JB01531-K-US
; CURRENT APPLICATION NUMBER: US/10/400,273
; CURRENT FILING DATE: 2003-03-26
; PRIOR APPLICATION NUMBER: 60/367,937
; PRIOR FILING DATE: 2002-03-27
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 411
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-400-273-1

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Query Match          100.0%; Score 144; DB 15; Length 411;
Best Local Similarity 100.0%; Pred. No. 4,1e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
    |||||
Db 29 GYVEMTVGSPPTLNIIVDTGSSNFAV 56

```

```

RESULT 12
US-10-627-473-19
; Sequence 19, Application US/10627473
; Publication No. US20040096950A1
; GENERAL INFORMATION:
; APPLICANT: VUILARD, LAURENT MICHEL MARIE
; APPLICANT: PATEL, SAHIL JOE
; APPLICANT: YON, JEFFREY ROLAND
; APPLICANT: CLAESBY, ANNE
; APPLICANT: HAMILTON, BRUCE JOHN

```

```

; APPLICANT: SHAH, ALEEM
; TITLE OF INVENTION: CRYSTAL STRUCTURE OF BETA SITE APP CLEAVING ENZYME
; FILE REFERENCE: 674553-2002.1
; CURRENT APPLICATION NUMBER: US/10/627,473
; CURRENT FILING DATE: 2003-07-25
; PRIOR APPLICATION NUMBER: 60/398,681
; PRIOR FILING DATE: 2002-07-26
; NUMBER OF SEQ ID NOS: 46
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 19
; LENGTH: 411
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-627-473-19

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Query Match          100.0%; Score 144; DB 15; Length 411;
Best Local Similarity 100.0%; Pred. No. 4,1e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
    |||||
Db 32 GYVEMTVGSPPTLNIIVDTGSSNFAV 59

```

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RESULT 13
US-10-627-473-20
; Sequence 20, Application US/10627473
; Publication No. US20040096950A1
; GENERAL INFORMATION:
; APPLICANT: VUILARD, LAURENT MICHEL MARIE
; APPLICANT: PATEL, SAHIL JOE
; APPLICANT: YON, JEFFREY ROLAND
; APPLICANT: CLAESBY, ANNE
; APPLICANT: HAMILTON, BRUCE JOHN
; APPLICANT: SHAH, ALEEM
; TITLE OF INVENTION: CRYSTAL STRUCTURE OF BETA SITE APP CLEAVING ENZYME
; FILE REFERENCE: 674553-2002.1
; CURRENT APPLICATION NUMBER: US/10/627,473
; CURRENT FILING DATE: 2003-07-25
; PRIOR APPLICATION NUMBER: 60/398,681
; PRIOR FILING DATE: 2002-07-26
; NUMBER OF SEQ ID NOS: 46
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 20
; LENGTH: 411
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-627-473-20

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Query Match          100.0%; Score 144; DB 15; Length 411;
Best Local Similarity 100.0%; Pred. No. 4,1e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
    |||||
Db 32 GYVEMTVGSPPTLNIIVDTGSSNFAV 59

```

```

RESULT 14
US-10-281-092-9
; Sequence 9, Application US/10281092
; Publication No. US20040121947A1
; GENERAL INFORMATION:
; APPLICANT: Ghosh, Arun K.
; APPLICANT: Tang, Jordan J.N.
; APPLICANT: Bilcer, Geoffrey
; APPLICANT: Chang, Manpin
; APPLICANT: Hong, Lin
; APPLICANT: Koelsch, Gerald E.
; APPLICANT: Loy, Jeffrey A.
; APPLICANT: Turner, Robert T., III

```

APPLICANT: Devasumadrum, Thippeswamy
TITLE OF INVENTION: COMPOUNDS WHICH INHIBIT BETA-SECRETASE
FILE REFERENCE: 2932.1001-004
CURRENT APPLICATION NUMBER: US/10/281,092
CURRENT FILING DATE: 2002-10-23
PRIOR APPLICATION NUMBER: US 10/032,818
PRIOR FILING DATE: 2001-12-28
PRIOR APPLICATION NUMBER: PCT US01/50826
PRIOR FILING DATE: 2001-12-28
PRIOR APPLICATION NUMBER: US 60/258,705
PRIOR FILING DATE: 2000-12-28
PRIOR APPLICATION NUMBER: US 60/275,756
PRIOR FILING DATE: 2001-03-14
PRIOR APPLICATION NUMBER: US 60/335,952
PRIOR FILING DATE: 2001-10-23
PRIOR APPLICATION NUMBER: US 60/333,545
PRIOR FILING DATE: 2001-11-27
PRIOR APPLICATION NUMBER: US 60/348,464
PRIOR FILING DATE: 2002-01-14
PRIOR APPLICATION NUMBER: US 60/348,615
PRIOR FILING DATE: 2002-01-14
PRIOR APPLICATION NUMBER: US 60/390,804
PRIOR FILING DATE: 2002-06-20
PRIOR APPLICATION NUMBER: US 60/397,557
PRIOR FILING DATE: 2002-07-19
Remaining Prior Application data removed - See File Wrapper or PALM.
NUMBER OF SEQ ID NOS: 59
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 9
LENGTH: 414
TYPE: PRT
ORGANISM: Unknown
FEATURE:
OTHER INFORMATION: memapsin 2
US-10-281-092-9

Query Match 100.0%; Score 144; DB 16; Length 414;
Best Local Similarity 100.0%; Pred. No. 4.1e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTNLIVDTGSSNPAV 28
DB 34 GYVEMTVGSPPTNLIVDTGSSNPAV 61

RESULT 15
US-10-627-473-21
Sequence 21, Application US/10627473
Publication No. US20040096950A1
GENERAL INFORMATION:
APPLICANT: VUILLARD, LAURENT MICHEL MARIE
APPLICANT: PATEL, SAHIL JOE
APPLICANT: VON, JEFFREY ROLAND
APPLICANT: CLEASBY, ANNE
APPLICANT: HAMILTON, BRUCE JOHN
APPLICANT: SHAH, ALEEM
TITLE OF INVENTION: CRYSTAL STRUCTURE OF BETA SITE APP CLEAVING ENZYME
FILE REFERENCE: 674553-2002.1
CURRENT APPLICATION NUMBER: US/10/627,473
CURRENT FILING DATE: 2003-07-25
PRIOR APPLICATION NUMBER: 60/398,681
PRIOR FILING DATE: 2002-07-26
NUMBER OF SEQ ID NOS: 46
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 21
LENGTH: 417
TYPE: PRT
ORGANISM: Homo sapiens
US-10-627-473-21

Query Match 100.0%; Score 144; DB 15; Length 417;

Best Local Similarity 100.0%; Pred. No. 4.2e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTNLIVDTGSSNPAV 28
DB 32 GYVEMTVGSPPTNLIVDTGSSNPAV 59

Search completed: July 26, 2005, 16:51:25
Job time : 154 secs

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OM protein - protein search, using sw model

Run on: July 26, 2005, 16:38:10 ; Search time 39 Seconds

(without alignments)
69.079 Million cell updates/sec

Title: US-10-726-967A-52

Perfect score: 144

Sequence: 1 GYVEMTVGSPPTINILVDTGSSNFAV 28

Scoring table: BLOSUM62
Gapop. 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 200000000Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 79:*

1: PIR1: *
2: PIR2: *
3: PIR3: *
4: PIR4: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	144	100.0	501	2 A59090	aspartic proteinase
2	91	63.2	709	2 T29692	hypothetical prote
3	90	62.5	396	2 T47207	aspartic proteinase
4	87	60.4	405	2 A25379	gastropepsin (EC
5	83	57.6	398	2 T33383	hypothetical prote
6	83	57.6	406	1 REHUK	renin (EC 3.4.23.1
7	82	56.9	383	2 UC7573	pepsinogen C - Alt
8	82	56.9	384	2 A39314	gastric (EC 3.4.2
9	82	56.9	389	2 JEO371	pepsin C (EC 3.4.2
10	82	56.9	400	2 I47099	renin (EC 3.4.23.1
11	82	56.9	401	1 REMSK	renin (EC 3.4.23.1
12	82	56.9	402	1 REMSK	renin (EC 3.4.23.1
13	82	56.9	506	2 S71591	aspartic proteinase
14	82	56.9	508	2 D85056	probable aspartic
15	82	56.9	509	2 S49349	cyprosin (EC 3.4.2
16	82	56.9	596	2 S57971	aspartic proteinase
17	81	56.2	387	2 A45117	hypothetical prote
18	81	56.2	433	2 E96649	aspartic prote
19	81	56.2	513	2 T09739	aspartic endopepti
20	81	56.2	537	2 S50344	hypothetical prote
21	80	55.6	506	2 F86253	probable aspartic
22	80	55.6	506	2 T07915	aspartic prote
23	80	55.6	375	2 S64957	aspartic prote
24	79	54.9	375	2 C96715	protein P4N2.8 (im
25	78	54.2	165	2 S61602	probable membrane
26	78	54.2	344	1 KHPGD	cathepsin D (EC 3.
27	78	54.2	377	1 PEMOCJ	gastric (EC 3.4.
28	78	54.2	388	2 JCT7246	pepsinogen C - com
29	78	54.2	388	2 A29937	gastric (EC 3.4

30	78	54.2	392	1 A24608	gastric (EC 3.4
31	78	54.2	398	2 I51185	cathepsin D (EC 3.
32	78	54.2	407	1 KHRUD	cathepsin D (EC 3.
33	78	54.2	412	1 KHRUD	cathepsin D (EC 3.
34	77	53.5	394	2 B43356	gastric (EC 3.4
35	76	52.8	391	2 A43356	cathepsin E (EC 3.
36	76	52.8	396	2 A34401	cathepsin E (EC 3.
37	76	52.8	508	2 S19697	aspartic proteinase
38	76	52.8	509	2 S66516	oryzasin (EC 3.4.2
39	75	52.1	383	2 A41443	pepsin (EC 3.4.23.
40	75	52.1	410	1 KXMSD	cathepsin D (EC 3.
41	75	52.1	438	2 S47096	cynarase (EC 3.4.2
42	75	52.1	425	2 T45035	hypothetical prote
43	75	52.1	474	2 T12049	cyprosin (EC 3.4.2
44	74	51.4	151	2 T18478	hypothetical prote
45	74	51.4	189	2 T18480	hypothetical prote

ALIGNMENTS

RESULT 1
A59090
aspartic proteinase (EC 3.4.23.-) BACE precursor - human
N:Alternate names: beta-secretase; beta-site APP cleaving enzyme
C:Species: Homo sapiens (man)
C>Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 09-Jul-2004
C:Accession: A59090
C:Vassar, R.; Bennett, B.D.; Babu-Khan, S.; Kahn, S.; Mendiaz, E.A.; Denis, P.; Teplow, M.A.; Biere, A.L.; Curran, E.; Burgess, T.; Louis, J.C.; Collins, F.; Treanor, J.; Roge, Science 286, 735-741, 1999
A>Title: beta-Secretase cleavage of Alzheimer's amyloid precursor protein by the transmembrane aspartic protease A59090; PMID:20002972; PMID:10531052
A:Reference number: A59090; PMID:20002972; PMID:10531052
A>Note: submitted to GenBank, September 1999
A:Accession: A59090
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-501 <VASS>
A:Cross-references: UNIPROT:P56817; GB:AF190725; NID:G6118538; PIDN:AAF04142.1; PID:G61
C:Genetics:
A:Gene: BACE
C:Superfamily: beta-secretase
C:Keywords: Alzheimer's disease; aspartic proteinase; brain; glycoprotein; hydrolase; P.
F:1-21/Domain: signal sequence #status predicted <SIG>
F:22-45/Domain: propeptide #status predicted <PRO>
F:46-501/Product: acid proteinase BACE #status predicted <MAT>
F:461-477/Domain: transmembrane #status predicted <TRN>
F:93,289/Active site: Asp #status predicted
F:153,172,223,354/Binding site: carbohydrate (asn) (covalent) #status predicted
F:330-380/Disulfide bonds: #status predicted

Query Match 100.0%; Score 144; DB 2; Length 501;
Best Local Similarity 100.0%; Pred. No. 7.3e-14;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTINILVDTGSSNFAV 28
DB 74 GYVEMTVGSPPTINILVDTGSSNFAV 101

RESULT 2
T29692
hypothetical protein T189.2 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
C:Accession: T29692
R:Du, Z.; Gatlund, S.
Submitted to the EMBL Data Library, November 1995
A:Description: The sequence of C. elegans cosmid T189.
A:Reference number: Z20666
A:Accession: T29692
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA

A;Residues: 1-709 <DUZ>
A;Cross-references: EMBL:U41746; PIDN:AAA63331.1; CESP:TI8H9.2
C;Genetics:
A;Gene: CESP:TI8H9.2
A;Introns: 4/3; 126/3; 161/3; 190/3; 239/3; 282/3; 401/1; 691/3

Query Match
Best Local Similarity 63.2%; Score 91; DB 2; Length 709;
Matches 16; Conservative 5; Mismatches 5; Indels 0; Gaps 0;

Db
1 GYVMTVGSPPQTLNLTVDGSSNF 26
39 GYVGTIVGSPQERRVMTDGSNF 64

RESULT 3
T47207
aspartic proteinase (EC 3.4.23.-) [imported] - Neurospora crassa
C;Species: Neurospora crassa
C;Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 09-Jul-2004
R;Bowman, B.
Submitted to the EMBL Data Library, September 1995
A;Reference number: Z24391
A;Accession: T47207
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-396 <BOW>
A;Cross-references: UNIPROT:Q01294; EMBL:U36471; PIDN:AAA79878.1
C;Genetics:
A;Gene: pep-4
A;Introns: 33/3; 85/1
C;Superfamily: pepsin
C;Keywords: aspartic proteinase; hydrolase

Query Match
Best Local Similarity 62.5%; Score 90; DB 2; Length 396;
Matches 15; Conservative 7; Mismatches 5; Indels 0; Gaps 0;

Db
2 YVEMTVGSPPQTLNLTVDGSSNF 28
85 YFSEITITGPQTFKVLVDGSSNLMV 111

RESULT 4
A25379
saccharopepsin (EC 3.4.23.25) precursor - yeast (Saccharomyces cerevisiae)
N;Alternate names: protein P2585; protein YPL154c; proteinase A; Saccharomyces aspartic
C;Species: Saccharomyces cerevisiae
C;Date: 25-Oct-1987 #sequence_revision 25-Oct-1987 #text_change 16-Aug-2004
C;Accession: A25379; A24711; B42640; S65165; S69445; S71635; S28231; S28233
R;Woolford, C.A.; Daniels, L.B.; Park, F.D.; Jones, E.W.; Van Ardeell, J.N.; Innis, M.A.
Mol. Cell. Biol. 6, 2500-2510, 1986
A;Title: The PEP4 gene encodes an aspartyl protease implicated in the posttranslational
A;Reference number: A25379; MUID:87064548; PMID:3537721
A;Accession: A25379
A;Molecule type: DNA
A;Residues: 1-405 <WOO>
A;Cross-references: UNIPROT:P07267; EMBL:M13358; NID:g172121; PIDN:AAB63975.1; PID:g1721
R;ammerer, G.; Hunter, C.P.; Rothman, J.H.; Saari, G.C.; Valls, L.A.; Stevens, T.H.
Mol. Cell. Biol. 6, 2490-2499, 1986
A;Title: PEP4 gene of Saccharomyces cerevisiae encodes proteinase A, a vacuolar enzyme
A;Reference number: A25378; MUID:87064547; PMID:3025936
A;Accession: A25378
A;Molecule type: DNA
A;Residues: 1-405 <AMM>
A;Cross-references: GB:M13358; GB:M13632; NID:g172121; PIDN:AAB63975.1; PID:g172122
R;Dwyer, T.; Halkier, B.; Svendsen, I.; Ottesen, M.
Carlsberg Res. Commun. 51, 27-41, 1986
A;Title: Primary structure of the aspartic proteinase A from Saccharomyces cerevisiae.
A;Reference number: A24711
A;Accession: A24711
A;Molecule type: protein

A;Residues: 77-405 <DRE>
R;Roof, D.M.; Meluh, P.B.; Rose, M.D.
J. Cell Biol. 118, 95-108, 1992
A;Title: Kinesin-related proteins required for assembly of the mitotic spindle.
A;Reference number: A42640; MUID:92317166; PMID:1618910
A;Accession: B42640
A;Molecule type: DNA
A;Residues: 373-405 <ROO>
A;Cross-references: EMBL:Z11963; NID:g3852; PIDN:CAA78020.1; PID:g3853
A;Note: the authors did not translate the codons for residues 373, 374, and 375
A;Note: sequence extracted from NCBI backbone (NCBI:107719, NCBI:107721)
R;Purnelle, B.; Coater, F.; Goffeau, A.
Submitted to the Protein Sequence Database, May 1996
A;Reference number: S65154
A;Accession: S65154
A;Molecule type: DNA
A;Residues: 1-405 <PUR>
A;Cross-references: EMBL:Z73510; NID:g1370327; PIDN:CAA97859.1; PID:g1370328; MIPS:YPL15
A;Experimental source: strain 5288C (AB972)
R;Purnelle, B.; Comblez, S.; Coater, F.; Naveau, F.; Goffeau, A.
Submitted to the EMBL Data Library, March 1996
A;Description: The sequence of 55 kb on the left arm of yeast chromosome XVI identifies
ogues to the human phosphotyrosyl phosphatase activator PRPA and a homologue to the plant
A;Reference number: S69448
A;Accession: S69448
A;Molecule type: DNA
A;Residues: 1-405 <PUM>
A;Cross-references: EMBL:X96770; NID:g1403537; PIDN:CAA65567.1; PID:g1403555
R;Wolff, A.M.; Din, N.; Little Peteresen, J.G.
Yeast 12, 823-832, 1996
A;Title: Vacuolar and extracellular maturation of Saccharomyces cerevisiae proteinase A
A;Reference number: S71635; MUID:56437971; PMID:8840499
A;Accession: S71635
A;Molecule type: protein
A;Residues: 23-31;68-86 <WOL>
C;Genetics:
A;Gene: SGD:PEP4; PRA1; PHO9
A;Cross-references: SGD:S0006075; MIPS:YPL154C
A;Map position: 16L
C;Function:
A;Description: responsible for degradation of internalized alpha-factor receptor and a-f
alkaline phosphatase, acid trehalase, and vacuolar exopolyphosphatase
C;Superfamily: Pepsin
C;Keywords: aspartic proteinase; hydrolase; yeast vacuole
F;1-22/Domain: signal sequence #status predicted <SIG>
F;23-67/Domain: propeptide #status experimental <PRO>
F;68-405/Product: saccharopepsin #status experimental <MAT>

Query Match
Best Local Similarity 60.4%; Score 87; DB 2; Length 405;
Matches 14; Conservative 7; Mismatches 6; Indels 0; Gaps 0;

Db
2 YVEMTVGSPPQTLNLTVDGSSNF 28
91 YVDITLITGPQTFKVLVDGSSNLMV 117

RESULT 5
T33383
hypothetical protein H22K11.1 - Caenorhabditis elegans
C;Species: Caenorhabditis elegans
C;Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 09-Jul-2004
R;Beck, C.; Wamsley, P.; Keppler, N.
Submitted to the EMBL Data Library, July 1998
A;Description: The sequence of C. elegans cosmid H22K11.
A;Reference number: Z21333
A;Accession: T33383
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-398 <BEC>
A;Cross-references: UNIPROT:P55956; EMBL:A077544; PIDN:AAC64617.1; GSPDB:GN00028; CESP:1
A;Experimental source: strain Bristol N2; clone H22K11

JC7573
 pepsinogen C - African clawed frog
 N:Alternate names: progastricsin
 C:Species: Xenopus laevis (African clawed frog)
 C>Date: 30-Jun-2001 #sequence_revision 30-Jun-2001 #text_change 09-Jul-2004
 C:Accession: JC7573; PC7118
 R:Ikuzawa, M.; Inokuchi, T.; Kobayashi, K.; Yasunasu, S.
 U:Biochem. 129, 147-153, 2001
 A:Title: Amphibian pepsinogens: Purification and characterization of Xenopus pepsinogens
 A:Reference number: JC7573; MUID:21064922; PMID:11134969
 A:Contents: Stomach
 A:Accession: JC7573
 A:Molecule type: mRNA
 A:Residues: 1-383 <IKU>
 A:Cross-references: UNIPROT:Q9DEC3; DDBJ:AB045379
 A:Accession: PC7118
 A:Molecule type: protein
 A:Residues: 17-68 <IK2>
 C:Comment: This protein is a zymogen for gastric aspartic proteinase, with pepsin-like
 C:Genetics:
 A:Gene: Pgc
 C:Superfamily: pepsin
 C:Keywords: stomach; zymogen

Query Match 56.9%; Score 82; DB 2; Length 383;
 Best Local Similarity 55.6%; Pred. No. 0.00015;
 Matches 15; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY 2 YVEMTVGSPQPTNLIVDTGSSNFAV 28
 Db 67 YVGEISIGTPQNFVLVFDTGSSNLWV 93

RESULT 8
 A39314
 gastricsin (EC 3.4.23.3) precursor - bullfrog
 C:Species: Rana catesbeiana (bullfrog)
 C>Date: 19-Jun-1992 #sequence_revision 19-Jun-1992 #text_change 16-Aug-2004
 C:Accession: A39314
 R:Yakabe, E.; Tanji, M.; Ichinose, M.; Goto, S.; Miki, K.; Kurokawa, K.; Ito, H.; Kageya
 J. Biol. Chem. 266, 22436-22443, 1991
 A:Title: Purification, characterization, and amino acid sequences of pepsinogens and pep
 A:Reference number: A39314; MUID:92042186; PMID:1939266
 A:Accession: A39314
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-384 <YAK>
 A:Cross-references: UNIPROT:Q91322; GB:M73750; NID:G213687; PIDN:AAA49530.1; PID:G21368
 C:Superfamily: Pepsin
 C:Keywords: aspartic proteinase; hydrolase; protein digestion

Query Match 56.9%; Score 82; DB 2; Length 384;
 Best Local Similarity 55.6%; Pred. No. 0.00015;
 Matches 15; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY 2 YVEMTVGSPQPTNLIVDTGSSNFAV 28
 Db 67 YVGEISIGTPQNFVLVFDTGSSNLWV 93

RESULT 9
 J50371
 pepsin C (EC 3.4.23.-) precursor - chicken
 N:Alternate names: pepsinogen C
 C:Species: Gallus gallus (chicken)
 C>Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 09-Jul-2004
 C:Accession: J50371
 R:Sakamoto, N.; Saiga, H.; Yasugi, S.
 Biochem. Biophys. Res. Commun. 250, 420-424, 1998
 A:Title: Analysis of temporal expression pattern and cis-regulatory sequences of chicken
 A:Reference number: J50370; MUID:98440813; PMID:9753645
 A:Accession: J50371
 A:Status: preliminary

A:Molecule type: mRNA
 A:Residues: 1-389 <SAK>
 A:Cross-references: UNIPROT:Q9W643; UNIPROT:Q9PWK1
 C:Superfamily: pepsin
 C:Keywords: aspartic proteinase; hydrolase

Query Match 56.9%; Score 82; DB 2; Length 389;
 Best Local Similarity 55.6%; Pred. No. 0.00016;
 Matches 15; Conservative 5; Mismatches 7; Indels 0; Gaps 0;

QY 2 YVEMTVGSPQPTNLIVDTGSSNFAV 28
 Db 73 YVGEISIGTPQNFVLVFDTGSSNLWV 99

RESULT 10
 I47099
 renin (EC 3.4.23.15) - sheep
 C:Species: Ovis orientalis aries, Ovis ammon aries (domestic sheep)
 C>Date: 15-Oct-1996 #sequence_revision 15-Oct-1996 #text_change 09-Jul-2004
 C:Accession: I47099
 R:Alfred, G.P.; Fu, P.; Crawford, R.J.; Fernley, R.T.
 J. Mol. Endocrinol. 8, 3-11, 1992
 A:Title: The sequence and tissue expression of ovine renin.
 A:Reference number: I47099; MUID:92181567; PMID:1543552
 A:Accession: I47099
 A:Status: preliminary; translated from GB/EMBL/DDBJ
 A:Molecule type: mRNA
 A:Residues: 1-400 <ALD>
 A:Cross-references: UNIPROT:P52115; GB:I43524; NID:G896317; PIDN:AAA69809.1; PID:G89631
 C:Superfamily: pepsin
 C:Keywords: aspartic proteinase; hydrolase
 F:98,286/Active site: Asp #status predicted

Query Match 56.9%; Score 82; DB 2; Length 400;
 Best Local Similarity 51.9%; Pred. No. 0.00016;
 Matches 14; Conservative 6; Mismatches 7; Indels 0; Gaps 0;

QY 2 YVEMTVGSPQPTNLIVDTGSSNFAV 28
 Db 80 YVGEISIGTPQTFKVIPTGSSANLWV 106

RESULT 11
 REMS
 renin (EC 3.4.23.15) precursor, submandibular - mouse
 C:Species: Mus musculus (house mouse)
 C>Date: 15-Oct-1982 #sequence_revision 17-Dec-1982 #text_change 16-Aug-2004
 C:Accession: A93283; A93285; B93285; B22058; A00988
 R:Mason, K.S.; Chang, J.U.; Inagami, T.
 Proc. Natl. Acad. Sci. U.S.A. 79, 4858-4862, 1982
 A:Title: Amino acid sequence of mouse submaxillary gland renin.
 A:Reference number: A93923; MUID:83014991; PMID:6812055
 A:Accession: A93923
 A:Molecule type: protein
 A:Residues: 64-351;354-401 <MIS>
 A:Cross-references: UNIPROT:P00796
 R:Pantlher, J.U.; Foote, S.; Chambrud, B.; Stroberg, A.D.; Corvol, P.; Rougeon, F.
 Nature 298, 90-92, 1982
 A:Title: Complete amino acid sequence and maturation of the mouse submaxillary gland ren
 A:Reference number: A93285; MUID:82220074; PMID:6283373
 A:Accession: A93285
 A:Molecule type: mRNA
 A:Residues: 1-98, 'W', 100-194, 'LSRS', 199-394, 'V', 396-401 <PA1>
 A:Cross-references: GB:J00621; GB:V00845; NID:G200701; PIDN:AAA40050.1; PID:G200702
 A:Note: the authors translated codon ATG for residue 99 as Ile
 A:Accession: B93285
 A:Molecule type: protein
 A:Residues: 64-84;354-374 <PA2>
 R:Poel, M.; Llesch, J.M.
 J. Biol. Chem. 258, 9856-9860, 1983
 A:Title: Mouse submaxillary gland renin contains a noncovalently attached fatty acid.
 A:Reference number: A92439; MUID:83290909; PMID:6350284

A/Contents: annotation; fatty acid binding
R/Pantlier, J.J.; Dreyfus, M.; Roux, D.T.L.; Rougeon, F.
Proc. Natl. Acad. Sci. U.S.A. 81, 5489-5493, 1984
A/Title: Mouse kidney and submaxillary gland renin genes differ in their 5' putative reg
A/Reference number: A22058; MUID:84298161; PMID:6089205
A/Accession: B22058
A/Molecule type: DNA
A/Residues: 1-29 <PAN>
C/Comment: The enzyme contains a noncovalently attached fatty acid.
C/Comment: Submandibular renin has catalytic and antigenic activities similar to renal r
C/Comment: This renin is synthesized in the submandibular gland of males only.
C/Genetics:
A/Gene: REN2
C/Superfamily: Pepsin
C/Keywords: aspartic proteinase; hydrolase; salivary gland; submandibular gland
F/1-21/Domain: signal sequence #status predicted <SIG>
F/22-63/Domain: activation peptide #status predicted <ACP>
F/64-351/Product: renin, submandibular, heavy chain #status experimental <RSH>
F/354-401/Product: renin, submandibular, light chain #status experimental <RSL>
F/101,286/Active site: Asp #status experimental
F/114-121,277-281,320-357/Diulfide bonds: #status predicted

Query Match 56.9%; Score 82; DB 1; Length 401;
Best Local Similarity 51.9%; Pred. No. 0.00016;
Matches 14; Conservative 6; Mismatches 7; Indels 0; Gaps 0;

QY 2 YVEMTVGSPPTNTLVDTGSSNFAY 28
DB 83 YVGEIGIGTPPQTFKVIPTDGSANLWV 109

RESULT 12
REMSK
renin (EC 3.4.23.15) precursor, renal - mouse
N/Alternate names: angiotensin-forming enzyme; angiotensinogenase; renin 1
C/Species: Mus musculus (house mouse)
C/Date: 30-Jun-1987 #sequence revision 30-Jun-1987 #text change 16-Aug-2004
C/Accession: A00989; S07636; A22766; A22058; I57576; A05137; JH0083
R/Holm, I.; Ollio, R.; Pantlier, J.J.; Rougeon, F.
EMBO J. 3, 557-562, 1984
A/Title: Evolution of aspartyl proteases by gene duplication: the mouse renin gene is on
A/Reference number: A00989; MUID:84182525; PMID:6370686
A/Accession: A00989
A/Molecule type: DNA
A/Residues: 1-402 <HOL>
A/Cross-references: UNIPROT:P06281; EMBL:X00850
R/Kim, W.S.; Murakami, K.; Nakayama, K.
Nucleic Acids Res. 17, 9480, 1989
A/Title: Nucleotide sequence of a cDNA coding for mouse Ren1 preprorenin.
A/Reference number: S07636; MUID:90067953; PMID:2685761
A/Accession: S07636
A/Molecule type: mRNA
A/Residues: 1-402 <KIM>
A/Cross-references: EMBL:X16642; NID:G53930; PIDN:CAA34636.1; PID:G53931
R/Mullins, U.J.; Butt, D.W.; Winders, J.D.; McTurk, P.; George, H.; Brammar, W.J.
EMBO J. 1, 1461-1466, 1982
A/Title: Molecular cloning of two distinct renin genes from the DBA/2 mouse.
A/Reference number: A90968; MUID:84207899; PMID:6327270
A/Accession: A22766
A/Molecule type: mRNA
A/Residues: 269-314, 'D', 316 <MUL>
R/Pantlier, J.J.; Dreyfus, M.; Roux, D.T.L.; Rougeon, F.
Proc. Natl. Acad. Sci. U.S.A. 81, 5489-5493, 1984
A/Title: Mouse kidney and submaxillary gland renin genes differ in their 5' putative reg
A/Reference number: A22058; MUID:84298161; PMID:6089205
A/Accession: A22058
A/Molecule type: DNA
A/Residues: 1-30 <PAN>
R/Field, L.J.; Philbrick, W.M.; Howles, P.N.; Dickinson, D.P.; McGowan, R.A.; Gross, K.W.
Mol. Cell. Biol. 4, 2321-2331, 1984
A/Title: Expression of tissue-specific Ren-1 and Ren-2 genes of mice: Comparative analys
A/Reference number: I57576; MUID:85085936; PMID:6392850
A/Accession: I57576

A/Status: preliminary; translated from GB/EMBL/DBU
A/Molecule type: DNA
A/Residues: 1-31 <RES>
A/Cross-references: GB:X02800; NID:9200689; PIDN:AAA40044.1; PID:G200690
C/Comment: The only known function of renal renin is to release angiotensin I from angio
creased sodium retention by the kidney.
C/Comment: Renal renin is synthesized by the juxtaglomerular cells of the kidney in resp
C/Genetics:
A/Gene: Ren-1
A/Introns: 31/2; 81/3; 123/1; 162/3; 228/2; 268/2; 316/3; 349/3
C/Superfamily: Pepsin
C/Keywords: aspartic proteinase; blood pressure control; glycoprotein; hydrolase; kidney
F/1-21/Domain: signal sequence #status predicted <SIG>
F/22-64/Domain: propeptide #status predicted <PRO>
F/65-402/Product: renin #status predicted <MAT>
F/59,139,320/Binding site: carbohydrate (Asn) (covalent) #status predicted
F/102,287/Active site: Asp #status predicted

Query Match 56.9%; Score 82; DB 1; Length 402;
Best Local Similarity 51.9%; Pred. No. 0.00016;
Matches 14; Conservative 6; Mismatches 7; Indels 0; Gaps 0;

QY 2 YVEMTVGSPPTNTLVDTGSSNFAY 28
DB 84 YVGEIGIGTPPQTFKVIPTDGSANLWV 110

RESULT 13
S71591
aspartic proteinase precursor, wound-induced - tomato
C/Species: Lycopersicon esculentum (tomato)
C/Date: 04-Feb-1998 #sequence revision 13-Feb-1998 #text change 09-Jul-2004
C/Accession: S71591
R/Schaller, A.; Ryan, C.A.
Plant Mol. Biol. 31, 1073-1077, 1996
A/Title: Molecular cloning of a tomato leaf cDNA encoding an aspartic protease, a system
A/Reference number: S71591; MUID:97000919; PMID:8843949
A/Accession: S71591
A/Status: preliminary
A/Molecule type: mRNA
A/Residues: 1-506 <SCH>
A/Cross-references: UNIPROT:Q40140; EMBL:L46681; NID:G951448; PIDN:AA18280.1; PID:G9514
C/Superfamily: oryzasin; sapsin repeat homology

Query Match 56.9%; Score 82; DB 2; Length 506;
Best Local Similarity 55.6%; Pred. No. 0.00021;
Matches 15; Conservative 4; Mismatches 8; Indels 0; Gaps 0;

QY 2 YVEMTVGSPPTNTLVDTGSSNFAY 28
DB 83 YVGEIGIGTPPQTFKVIPTDGSANLWV 109

RESULT 14
D85056
probable aspartic proteinase [imported] - Arabidopsis thaliana
C/Species: Arabidopsis thaliana (mouse-ear cress)
C/Date: 16-Feb-2001 #sequence revision 16-Feb-2001 #text change 09-Jul-2004
C/Accession: D85056
R/anonymous, The European Union Arabidopsis Genome Sequencing Consortium, The Cold Sprin
Nature 402, 769-777, 1999
A/Title: Sequence and analysis of chromosome 4 of the plant Arabidopsis thaliana.
A/Reference number: A85001; MUID:20083408; PMID:10617198
A/Accession: D85056
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-508 <STO>
A/Cross-references: UNIPROT:Q9XEC4; GB:NC_001268; NID:G7267203; PIDN:CAB77914.1; GSPDB:Q
C/Genetics:
A/Gene: AtPg04460
A/Map position: 4
C/Superfamily: oryzasin; sapsin repeat homology

Query Match 56.9%; Score 82; DB 2; Length 508;
 Best Local Similarity 54.2%; Pred. No. 0.00021;
 Matches 13; Conservative 6; Mismatches 5; Indels 0; Gaps 0;

OY 2 YVEMTVGSPQPTLNILVDTGSSN 25
 ||:::|||||:::|||||
 Db 87 YGDIITIGTPQKFTVIFDTGSSN 110

RESULT 15

S49349
 Cyprosin (EC 3.4.23.-) - cardoon
 C:Species: Cynara cardunculus (cardoon)
 C>Date: 16-Feb-1995 #sequence_revision 12-May-1995 #text_change 09-Jul-2004
 C:Accession: S49349
 R:Pietrzak, M.; Brodelius, P.; Pais, M.S.
 submitted to the EMBL Data Library, September 1994
 A:Reference number: S49349
 A:Accession: S49349
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-509 <PIB>
 A:Cross-references: UNIPROT:Q39476; EMBL:X81984; NID:9556818; PIDN:CA57510.1; PID:95568
 C:Comment: The pair of saposin repeat homology domains tagged SAP1 and SAP2 represent a
 C:Superfamily: oryzasin; saposin repeat homology
 C:Keywords: aspartic proteinase; hydrolase
 F:316-361/Domain: saposin repeat homology #status atypical <SAP1>
 F:375-420/Domain: saposin repeat homology #status atypical <SAP2>
 F:103,290/Active site: Asp #status predicted

Query Match 56.9%; Score 82; DB 2; Length 509;
 Best Local Similarity 55.6%; Pred. No. 0.00021;
 Matches 15; Conservative 4; Mismatches 8; Indels 0; Gaps 0;

OY 2 YVEMTVGSPQPTLNILVDTGSSNFAV 28
 ||:::|||||:::|||||
 Db 85 YGDIITIGTPQKFTVIFDTGSSNLMV 111

Search completed: July 26, 2005, 16:47:58
 Job time : 40 secs

GenCore version 5.1.6
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OM proteoin - protein search, using sw model

Run on: July 26, 2005, 16:37:25 ; Search time 167 Seconds

(without alignments)
85.858 Million cell updates/sec

Title: US-10-726-967A-52

Perfect score: 144

Sequence: 1 GYVEMTVGSPQTLNLTVDTGSSNPAV 28

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1612378 seqs, 512079187 residues

Total number of hits satisfying chosen parameters: 1612378

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

Database : UniProt 03:.*
1: uniprot_sprot:.*
2: uniprot_trembl:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	144	100.0	127	2	076KP0
2	144	100.0	457	2	08CAF4
3	144	100.0	501	1	BAE1_HUMAN
4	144	100.0	501	1	BAE1_MOUSE
5	144	100.0	501	1	BAE1_RAT
6	144	100.0	501	2	081YC8
7	144	100.0	501	2	08BOY4
8	144	100.0	532	2	09DUS1
9	143	99.3	501	2	08CTRI
10	124	86.1	505	2	06N2T7
11	122	84.7	500	2	07T0Y2
12	120	83.3	499	2	06PB20
13	119	82.6	396	2	09NZL1
14	119	82.6	439	2	09H2V8
15	119	82.6	468	2	09NZL2
16	119	82.6	518	1	BAE2_HUMAN
17	116	80.6	514	2	06IE75
18	116	80.6	514	2	08CSE9
19	116	80.6	514	2	08C793
20	116	80.6	514	2	09JUL8
21	100	69.4	244	2	08WQJ9
22	93	64.6	423	2	08N2D4
23	91	63.2	396	2	06C080
24	90	62.5	396	1	CARP_NEUCR
25	90	62.5	396	2	07RVB0
26	90	62.5	409	2	06CRM3
27	89	61.8	173	2	096TV7
28	89	61.8	427	2	P91802
29	89	61.8	428	2	07JNB4
30	89	61.8	429	2	026515
31	87	60.4	405	1	CARP_YEAST

32	86	59.7	398	2	06H2K5	06H2K5 botrytis ci
33	86	59.7	400	2	086ZP8	086ZP8 paracoccidi
34	86	59.7	504	2	093XR0	093XR0 ipomoea bat
35	85	59.0	354	2	09GYX7	09GYX7 boophilius m
36	84	58.3	397	2	08NUS2	08NUS2 leptosphaer
37	84	58.3	401	2	06TMM6	06TMM6 xenopus lae
38	84	58.3	406	2	06DLW5	06DLW5 macaca muli
39	83	57.6	358	2	09FRW5	09FRW5 nepenthes a
40	83	57.6	398	1	ASP3_CAEEL	P55956 caenorhabdi
41	83	57.6	400	1	RENI_CALJA	094522 callithrix
42	83	57.6	403	2	06TSC2	06TSC2 homo sapien
43	83	57.6	406	1	RENI_HUMAN	P00797 homo sapien
44	83	57.6	406	1	RENI_PANTR	P60016 pan troglod
45	83	57.6	406	2	06DLJ0	06DLJ0 macaca fasc

ALIGNMENTS

RESULT 1
076KP0 PRELIMINARY; PRT; 127 AA.
AC 076KP0;
DT 05-JUL-2004 (TRMBLrel. 27, Created)
DT 05-JUL-2004 (TRMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TRMBLrel. 27, Last annotation update)
DE Beta-site APP cleaving enzyme isoform I-127.
GN Name=BACE;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_Taxid=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Tanahashi H.;
RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.
CC 1-SIMILARITY: Belongs to peptidase family A1.
DR EMBL; AB089958; BAC81826.1; -
DR HSSP; P00797; 1BBS.
DR GO; GO:0009049; F:aspartic-type signal peptidase activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR001461; Peptidase_A1.
DR InterPro; IPR009119; Pept_A1_BACE.
DR InterPro; IPR009120; Pept_A1_BACE.
DR InterPro; IPR009007; Pept_Aspartic.
DR InterPro; IPR001969; Pept_Asp_AS.
DR Pfam; PF00026; Asp_1.
DR PRINTS; PR01816; BACEF.
DR PRINTS; PR01815; BACEFAMILY.
DR PROSITE; PS00141; ASP_PROTEASE; 1.
KW Aspartyl protease; Hydrolase; Protease.
SQ SSQENCE 127 AA; 13939 MW; C657354CBER72DC4 CRC64;
Query Match 100.0%; Score 144; DB 2; Length 127;
Best Local Similarity 100.0%; Pred. No. 4.6e-14;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GYVEMTVGSPQTLNLTVDTGSSNPAV 28
Db 74 GYVEMTVGSPQTLNLTVDTGSSNPAV 101
RESULT 2
08CAF4 PRELIMINARY; PRT; 467 AA.
AC 08CAF4;
DT 01-MAR-2003 (TRMBLrel. 23, Created)
DT 01-MAR-2003 (TRMBLrel. 23, Last sequence update)
DT 01-MAR-2004 (TRMBLrel. 26, Last annotation update)
DE Mus musculus 0 day neonate cerebellum cDNA, RIKEN full-length enriched
DE library, clone: C230037E16 product: beta-site APP cleaving enzyme, full
DE insert sequence.
GN Name=Bace1; Synonyms=Bace;

OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 ON NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=99279253; PubMed=10349636; DOI=10.1016/S0076-6879(99)03004-9;
 RA Carninci P., Hayashizaki Y.;
 RT "High-efficiency full-length cDNA cloning.";
 RL Meth. Enzymol. 303:19-44(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=21085660; PubMed=11217851; DOI=10.1038/35055500;
 RA RIKEN FANTOM Consortium;
 RT "Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 409:665-690(2001).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RA The FANTOM Consortium;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 Nature 420:563-573(2002).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=20499374; PubMed=11042159; DOI=10.1101/gr.145100;
 RA Carninci P., Shibata Y., Hayatsu N., Sugahara Y., Shibata K., Itoh M.,
 Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
 RT "Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new genes.";
 RL Genome Res. 10:1617-1630(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RX MEDLINE=20530913; PubMed=11076861; DOI=10.1101/gr.152600;
 RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
 Kono H., Akiyama J., Nishi K., Katsunai T., Tashiro H., Itoh M.,
 Sumi N., Ishii Y., Nakamura S., Hazama M., Nishino T., Harada A.,
 Yamamoto R., Matsunoto H., Sakaguchi S., Ikegami T., Kashiwagi K.,
 Fujiwaka S., Inoue K., Togawa Y., Izawa M., Ohata E., Wachihi K.,
 Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsura S., Kawai J.,
 Okazaki Y., Muramatsu M., Inoue Y., Kira A., Hayashizaki Y.;
 RT "RIKEN integrated sequence analysis (RISA) system-384-format
 sequencing pipeline with 384 multicapillary sequencer.";
 RL Genome Res. 10:1757-1771(2000).
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
 RA Adachi J., Aizawa K., Akimura T., Arakawa T., Bono H., Carninci P.,
 Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
 Hayashida K., Hayatsu N., Hiramoto K., Hiraoka T., Hirozane T.,
 Hori F., Imotani K., Ishii Y., Itoh M., Kagawa I., Kanakura T.,
 Katoh H., Kawai J., Kojima Y., Kondo S., Kono H., Kouda M., Koya S.,
 Kurihara C., Matsuyama T., Miyazaki A., Murata M., Nakamura M.,
 Nishi K., Nomura K., Numazaki R., Ohno M., Ohsato N., Okazaki Y.,
 Saio R., Saion H., Sakai C., Sakai K., Sakazume N., Sano H.,
 Sasaki D., Shibata K., Shingawa A., Shiraki T., Sogabe Y., Tagami M.,
 Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
 Tomaru A., Toya T., Yasunishi A., Muramatsu M., Hayashizaki Y.;
 RL Submitted (Apr-2002) to the EMBL/GenBank/DBJ databases.
 CC -1 - SIMILARITY: Belongs to peptidase family A1.
 DR EMBL: AK082317; BAC38462.1; -;
 DR HSSP: P56817; IPRN.
 DR MGD: MGI:1346542; Bace1.
 DR GO: GO:0005768; C:cytosol; ISS.
 DR GO: GO:0005615; C:extracellular space; TAS.
 DR GO: GO:0005794; C:Golgi apparatus; ISS.
 DR GO: GO:0016021; C:integral to membrane; ISS.
 DR GO: GO:0004190; F:aspartic-type endopeptidase activity; ISS.

DR GO: GO:0050435; P:beta-amyloid metabolism; ISS.
 DR GO: GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro: IPR001461; Peptidase A1.
 DR InterPro: IPR009119; Pept_A1_BACE.
 DR InterPro: IPR009120; Pept_A1_BACE1.
 DR InterPro: IPR009007; Pept_Asp_AS.
 DR InterPro: IPR001969; Pept_Asp_AS.
 DR PRINTS: PRO1816; BACE1.
 DR PRINTS: PRO1815; BACEFAMILY.
 DR PRINTS: PRO0792; PEPsin.
 DR PROSITE: PS00141; ASP_PROTEASE_1.
 DR ASpartyl protease; Hydrolase; Protease.
 SQ SEQUENCE 467 AA; 52063 MW; 31AB674FF1843652 CRC64;
 Query Match 100.0%; Score 144; DB 2; Length 467;
 Best Local Similarity 100.0%; Pred. No. 2, 2e-13;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GYVEMTVGSPQTLNLTDTGSSNFAV 28
 DB 74 GYVEMTVGSPQTLNLTDTGSSNFAV 101
 RESULT 3
 BAE1_HUMAN STANDARD; PRT; 501 AA.
 AC P56817; Q9BYB9; Q9BYC0; Q9BYC1; Q9BYT5;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 25-OCT-2004 (Rel. 45, Last annotation update)
 DE Beta-secretase 1 precursor (BC 3.4.23.46) (Beta-site APP cleaving
 enzyme 1) (Beta-site amyloid precursor protein cleaving enzyme 1)
 DE (Aspartyl protease 2) (Asp 2) (ASP2) (Membrane-associated aspartic
 protease 2) (Memapsin-2).
 GN Name=BACE1; Synonyms=BACE;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 ON NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORM A).
 RC TISSUE=Brain;
 RX MEDLINE=20002972; PubMed=10531052; DOI=10.1126/science.286.5440.735;
 RA Vaasat R., Bennett B.D., Babu-Khan S., Kahn S., Mendiaz E.A.,
 Denis P., Teplow D.B., Ross S., Amarante P., Loebner R., Luo Y.,
 Fisher S., Fuller J., Edenson S., Lile J., Jarosinski M.A.,
 Biere A.L., Curran E., Burgess T., Louis J.C., Collins F.,
 Treanor J., Rogers G., Citron M.;
 RT "Beta-secretase cleavage of Alzheimer's amyloid precursor protein by
 the transmembrane aspartic protease BACE.";
 RL Science 286:735-741(1999).
 RN [2]
 RP SEQUENCE FROM N.A. (ISOFORM A), SEQUENCE OF 46-68, AND
 RC CHARACTERIZATION.
 RP TISSUE=Brain;
 RX MEDLINE=20057171; PubMed=10591214; DOI=10.1038/990114;
 RA Sinha S., Anderson J.P., Barbour R., Basl G.S., Caccavello R.,
 Davis D., Dean M., Dovey H.F., Frigon N., Hong J., Jacobson-Croak K.,
 Jewett N., Keim P., Knops J., Lieberburg I., Fowler M., Tan H.,
 Tetsuno G., Tung J., Schenk D., Seubert P., Suenemasa S.M., Wang S.,
 Walker D., Zhao J., McConlogue L., Varghese J.;
 RT "Purification and cloning of amyloid precursor protein beta-secretase
 from human brain.";
 RL Nature 402:537-540(1999).
 RN [3]
 RP SEQUENCE FROM N.A. (ISOFORM A)
 RX MEDLINE=20057170; PubMed=10591213; DOI=10.1038/990107;
 RA Van R., Bienkowski M.J., Snuck M.E., Miao H., Tori M.C., Paulley A.M.,
 Braehler J.R., Strachan N.C., Mathews W.R., Buhl A.E., Carter D.B.,
 Tomasselli A.G., Parodi L.A., Heinrichson R.L., Guney M.E.;
 RT "Membrane-anchored aspartyl protease with Alzheimer's disease beta-
 secretase activity.";
 RL Nature 402:533-537(1999).

[4] SEQUENCE FROM N.A. (ISOFORM A).
 MEDLINE:20120043; PubMed:10656250; DOI=10.1006/mcne.1999.0811;
 RA Hussain I., Powell D.J., Howlett D.R., Tew D.G., Meek T.D.,
 RA Chapman C., Gloger I.S., Murphy K.B., Southan C.D., Ryan D.M.,
 RA Smith T.S., Simmons D.L., Walsh F.S., Dingwall C., Christie G.;
 RT "Identification of a novel aspartic proteinase (Asp 2) as beta-
 RT secretase";
 RL Mol. Cell. Neurosci. 14:419-427(1999).
 [5] SEQUENCE FROM N.A. (ISOFORM B).
 RC TISSUE=Brain, and Pancreas;
 RA Michel B., De Pietri Tonelli D., Zaccchetti D., Keller P.;
 RT "New beta-site APP cleaving enzyme isoform (BACE-1B) obtained from
 RT human brain and pancreas";
 RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
 [6] SEQUENCE FROM N.A. (ISOFORM C).
 RC TISSUE=Pancreas;
 RA Zaccchetti D., De Pietri Tonelli D., Schnurbe R.;
 RT "New beta-site APP cleaving enzyme isoform (BACE-1C) obtained from
 RT human pancreas";
 RL Submitted (JAN-2001) to the EMBL/GenBank/DBJ databases.
 [7] SEQUENCE FROM N.A. (ISOFORMS B; C AND D).
 RC TISSUE=Brain;
 RA MEDLINE:21408467; PubMed:11516562; DOI=10.1016/S0304-3940(01)01912-7;
 RA Tanahashi H., Tabira T.;
 RT "Three novel alternatively spliced isoforms of the human beta-site
 RT amyloid precursor protein cleaving enzyme (BACE) and their effect on
 RT amyloid beta-peptide production";
 RL Neurosci. Lett. 307:9-12(2001).
 [8] SEQUENCE OF 14-501 FROM N.A. (ISOFORM A), AND CHARACTERIZATION.
 RP MEDLINE:20144060; PubMed:10677483; DOI=10.1073/pnas.97.4.1456;
 RA Lin X., Koelsch G., Wu S., Downs D., Dashi A., Tang J.;
 RT "Human aspartic protease memapsin 2 cleaves the beta-secretase site of
 RT beta-amyloid precursor protein";
 RL Proc. Natl. Acad. Sci. U.S.A. 97:1456-1460(2000).
 [9] DISULFIDE BONDS.
 RP MEDLINE:21950860; PubMed:11953458;
 RA Fischer F., Molinari M., Bodendorf U., Paganetti P.;
 RT "The disulphide bonds in the catalytic domain of BACE are critical but
 RT not essential for amyloid precursor protein processing activity";
 RL J. Neurochem. 80:1079-1088(2002).
 -1- FUNCTION: Responsible for the proteolytic processing of the
 CC amyloid precursor protein (APP). Cleaves at the amino terminus of
 CC the A-beta peptide sequence, between residues 671 and 672 of APP,
 CC leads to the generation and extracellular release of beta-cleaved
 CC soluble APP, and a corresponding cell-associated carboxy-terminal
 CC fragment which is later released by gamma-secretase.
 -1- CATALYTIC ACTIVITY: Broad endopeptidase specificity. Cleaves Glu-
 CC Val-Asn-Ileu-| Asp-Ala-Glu-Phe in the Swedish variant of
 CC Alzheimer's amyloid precursor protein.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=4;
 CC Name=A; Synonyms=BACE-1A, BAC-501;
 CC IsoId=P56817-1; Sequence=Displayed;
 CC Name=B; Synonyms=BACE-1B, BACE-I-476;
 CC IsoId=P56817-2; Sequence=VSP_005223;
 CC Name=C; Synonyms=BACE-1C, BACE-I-457;
 CC IsoId=P56817-3; Sequence=VSP_005222;
 CC Name=D; Synonyms=BACE-1D, BACE-I-432;
 CC IsoId=P56817-4; Sequence=VSP_005222, VSP_005223;
 CC -1- TISSUE SPECIFICITY: Brain.
 CC -1- SIMILARITY: Belongs to the peptidase A1 family.
 CC -----
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 CC or send an email to license@sib-sib.ch).
 CC -----
 CC EMBL; AF190725; AAF04142.1; -
 DR EMBL; AF201468; AAF18982.1; -
 DR EMBL; AF200343; AAF17079.1; -
 DR EMBL; AF204943; AAF26367.1; -
 DR EMBL; AF338816; AAK38374.1; -
 DR EMBL; AF338817; AAK38375.1; -
 DR EMBL; AB050436; BAB40931.1; -
 DR EMBL; AB050437; BAB40932.1; -
 DR EMBL; AB050438; BAB40933.1; -
 DR EMBL; AF200193; AAF13715.1; -
 DR PIR; A59090; A59090.
 DR PDB; 1FXN; X-ray; A/B=56-446.
 DR PDB; 1M4H; X-ray; A/B=56-446.
 DR MEROPS; A01.004; -
 DR Genew; HGNC:933; BACE1.
 DR H-InvDB; HIX0010165; -
 DR MIM; 604252; -
 DR GO; GO:0005887; C:integral to plasma membrane; TAS.
 DR GO; GO:0005798; F:beta-aspartyl-peptidase activity; TAS.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; TAS.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009120; Pept_A1_BACE1.
 DR InterPro; IPR009169; Pept_Asp_AS.
 DR InterPro; IPR009007; Pept_Aspartic.
 DR InterPro; IPR001461; Peptidase_A1.
 DR Pfam; PF00026; Asp; 1.
 DR PRINTS; PRO1816; BACE1.
 DR PRINTS; PRO1815; BACEFAMILY.
 DR PRINTS; PRO0792; BPSIN.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.
 DR 3D-structure; Alternative splicing; Aspartyl protease;
 KW Direct protein sequencing; Glycoprotein; Hydrolase; Signal;
 KW Transmembrane; Zymogen.
 FT PROPEP 1 21
 FT SIGNAL 22 45
 FT CHAIN 46 501
 FT DOMAIN 22 457
 FT TRANSMEM 458 478
 FT DOMAIN 479 501
 FT ACT_SITE 93 93
 FT ACT_SITE 289 289
 FT DISULFID 216 420
 FT DISULFID 278 443
 FT DISULFID 330 380
 FT CARBOHYD 153 153
 FT CARBOHYD 172 172
 FT CARBOHYD 223 223
 FT CARBOHYD 354 354
 FT VARSPPLIC 146 189
 FT VARSPPLIC 190 214
 FT HELIX 61 63
 FT TURN 64 65
 FT STRAND 67 70
 FT TURN 71 73
 FT STRAND 74 81
 FT TURN 82 85
 FT STRAND 86 93
 FT TURN 94 95
 FT STRAND 99 102
 FT TURN 107 108
 FT HELIX 115 117
 FT TURN 119 120
 FT STRAND 122 131
 FT STRAND 136 147
 FT TURN 149 150
 FT STRAND 155 167
 FT TURN 172 173
 N-linked (GlcNAc. . .) (Potential).
 N-linked (GlcNAc. . .) (Potential).
 N-linked (GlcNAc. . .) (Potential).
 N-linked (GlcNAc. . .) (Potential).
 N-linked (in isoform C and isoform D).
 Missing (in isoform C and isoform D).
 Missing (in isoform B and isoform D).
 /FTId=VSP_005222.
 /FTId=VSP_005223.

FT STRAND 178 181
 FT HELIX 185 187
 FT TURN 192 193
 FT HELIX 197 204
 FT STRAND 211 215
 FT HELIX 224 229

Query Match 100.0%; Score 144; DB 1; Length 501;
 Best Local Similarity 100.0%; Pred. No. 2,4e-13;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTQNTINIVDTGSSNFAY 28
 DB 74 GYVEMTVGSPPTQNTINIVDTGSSNFAY 101

RESULT 4
 BAE1_MOUSE STANDARD; PRT; 501 AA.

AC 30-MAY-2000 (Rel. 39, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 25-OCT-2004 (Rel. 45, Last annotation update)
 DE Beta-secretase 1 precursor (BC 3.4.23.46) (Beta-site APP cleaving enzyme 1) (Beta-site amyloid precursor protein cleaving enzyme 1)
 DE (Aspartyl) protease 2) (ASP 2) (Membrane-associated aspartic protease 2) (Memapsin-2).
 GN Name=Bace1; Synonyms=Bace;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 CX NCBI_TaxID=10099;
 [1]

RP SEQUENCE FROM N.A.
 RX MEDLINE=20002972; PubMed=10531052; DOI=10.1126/science.286.5440.735;
 RA Vassar R., Bennett B.D., Babu-Khan S., Kahn S., Mendiaz E.A., Denis P., Teplow D.B., Ross S., Amarante P., Loebner R., Luo Y., Fisher S., Fuller J., Edenson S., Lile J., Jeromin M.A., Biere A.L., Curran E., Burgess T., Louis J.C., Collins F., Treanor J., Rogers G., Citron M.;
 RA "beta-secretase cleavage of Alzheimer's amyloid precursor protein by the transmembrane aspartic protease BACE";
 RL Science 286:735-741(1999).
 [2]

RP REVISIONS TO 6 AND 81-87.
 RL Bennett B.D., Vassar R., Citron M.;
 RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
 [3]

RP SEQUENCE FROM N.A.
 RX MEDLINE=20057170; PubMed=10591213; DOI=10.1038/990107;
 RA Yan R., Bienkowski M.J., Shuck M.E., Miao H., Tory M.C., Pauley A.M., Braheier J.R., Stratan N.C., Mathews W.R., Buhl A.E., Carter D.B., Tomasselli A.G., Parodi L.A., Heinrikson R.L., Gurney W.B.;
 RA "Membrane-anchored aspartyl protease with Alzheimer's disease beta-secretase activity";
 RL Nature 402:533-537(1999).
 [4]

RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Head;
 RX MEDLINE=222554683; PubMed=12466651; DOI=10.1038/nature01266;
 RA Okazaki Y., Furuno M., Kasukawa T., Adachi J., Bono H., Kondo S., Nikaido I., Oatso N., Saito R., Suzuki H., Yamana H., Kiyosawa H., Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Schonbach C., Gojohori T., Baldairelli R., Hill D.P., Bult C., Hume D.A., Quackenbush J., Schiml L.M., Kanapin A., Matsumoto S., Beisel K.W., Blake J.A., Brad D., Bruneic V., Chochia C., Corbani L.E., Cousins S., Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Frazer K.S., Grimmond S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D., Kanai A., Kawai H., Kawasawa Y., Kedzierski R.M., King B.L., Konagaya A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A., Maglocz D.R., Maltas L., Marchionni L., McKenzie L., Miki H., Nagashima T., Numata K., Okido T., Pavan W.J., Pertea G., Pesole G.,

RA Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S., Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M., Sandelin A., Schneider C., Sempke C.A., Setou M., Shimada K., Sultana R., Takenaka Y., Taylor M.S., Teasdale R.D., Tomita M., Verardo R., Wagner L., Wahlestedt C., Wang Y., Watanabe Y., Wells C., Wilming L.G., Wymshew-Boris A., Yanagisawa M., Yang I., Yang L., Yuan Z., Zavoian M., Zhu Y., Zimmer A., Carlini P., Hayatsu N., Hirozane-Kishikawa T., Kono H., Nakamura M., Sakazume N., Sato K., Shiroki T., Waki K., Kawai J., Aizawa K., Arakawa T., Fukuda S., Hara A., Hashizume W., Imotani K., Ishii Y., Itoh M., Kagawa I., Miyazaki A., Sakai K., Sasaki D., Shibata K., Shinagawa A., Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J., Birney E., Hayashizaki Y.;
 RA "Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs";
 RL Nature 420:563-573(2002).
 [5]

RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Brain;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G., Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K., Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F., Diatchenko L., Marusina K., Farmer A.A., Rubin G.W., Hong L., Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Ueda T.B., Tomihataki S., Carninci P., Prange C., Rhee S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullen S.J., Bosak S.A., McEwen P.J., McKernan K.J., Melek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hultky S.W., Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A., Fahey J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A., Whitting M., Madan A., Young A.C., Snevchenko Y., Bouffard G.G., Blakeley K.W., Touchman J.W., Green E.D., Dickson M.C., Rodriguez A.C., Grimwood J., Schultz J., Myers R.M., Butcherfield Y.S.N., Krzyzanski M.I., Skalska U., Smailus D.E., Schermer A., Schein J.E., Jones S.J.M., Marra M.A.;
 RA "Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 [6]

CC -I- FUNCTION: Responsible for the proteolytic processing of the amyloid precursor protein (APP). Cleaves at the amino terminus of the A-beta peptide sequence, between residues 671 and 672 of APP, leads to the generation and extracellular release of beta-cleaved soluble APP, and a corresponding cell-associated carboxy-terminal fragment which is later released by gamma-secretase (by similarity).
 CC -I- CATALYTIC ACTIVITY: Broad endopeptidase specificity. Cleaves Glu-Val-Asn-Leu-I-Asp-Ala-Glu-Phe in the Swedish variant of Alzheimer's amyloid precursor protein.
 CC -I- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -I- TISSUE SPECIFICITY: Brain.
 CC -I- SIMILARITY: Belongs to the peptidase A1 family.

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CC EMBL: AF190726; AAF04143.2;
 CC EMBL: AF200346; AAF17082.1;
 CC EMBL: AK014464; BAB29370.1;
 CC EMBL: BC048189; AAH48189.1;
 CC HSSP: P56817; 1MAH.
 CC MOPROS: A01.004;
 CC MGD: MG1:1346542; Bace1.
 CC InterPro: IPR009119; Pept_A1_BACE.
 CC InterPro: IPR009120; Pept_A1_BACE1.
 CC InterPro: IPR001969; Pept_Asp_AS.
 CC InterPro: IPR009007; Pept_AspArtic.

DR InterPro: IPR001461; Peptidase_A1.
DR Pfam: PF00026; Asp. 1.
DR PRINTS; PRO1816; BACE1.
DR PRINTS; PRO1815; BACEFAMILY.
DR PRINTS; PRO0792; PEPsin.
DR PROSITE; PS00141; ASP_PROTEASE; 1.
KW Aspartyl protease; Glycoprotein; Hydrolase; Signal; Transmembrane;
KM Zymogen.
FT SIGNAL 1 21 Potential.
FT PROPEP 22 45 Potential.
FT CHAIN 46 501 Beta-secretase 1.
FT DOMAIN 22 457 Extracellular (Potential).
FT TRANSMEM 458 478 Potential.
FT DOMAIN 479 501 Cytoplasmic (Potential).
FT ACT_SITE 93 93 By similarity.
FT ACT_SITE 289 289 By similarity.
FT DISULFID 216 420 By similarity.
FT DISULFID 278 443 By similarity.
FT DISULFID 330 380 By similarity.
FT CARBOHYD 153 153 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 172 172 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 223 223 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 354 354 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 501 AA; 55747 MW; C085A013145E474E CRC64;

Query Match 100.0%; Score 144; DB 1; Length 501;
Best Local Similarity 100.0%; Pred. No. 2,4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTQILNIVDTGSSNPAV 28
Db 74 GYVEMTVGSPPTQILNIVDTGSSNPAV 101
|||||
RESULT 5
ID BAE1_RAT STANDARD; PRT; 501 AA.
AC P56819;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 25-OCT-2004 (Rel. 45, Last annotation update)
DE Beta-secretase 1 precursor (EC 3.4.23.46) (Beta-site APP cleaving
enzyme 1) (beta-site amyloid precursor protein cleaving enzyme 1)
DE (Aspartyl) protease 2) (Asp 2) (ASP2) (Membrane-associated aspartic
protease 2) (Memapsin-2).
GN Name=BACE1; Synonyms=BACE;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Scturognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20002972; Pubmed=10531052; DOI=10.1126/science.286.5440.735;
RA Vassar R., Bennett B.D., Babu-Khan S., Kahn S., Mendiaz E.A.,
RA Denis P., Teplow D.B., Ross S., Amarante P., Loebner R., Luo Y.,
RA Fisher S., Fuller J., Edenson S., Lile J., Jarosinski M.A.,
RA Biere A.L., Curran E., Burgess T., Louis J.-C., Collins F.,
RA Treanor J., Rogers G., Citron M.,
RA "Beta-secretase cleavage of Alzheimer's amyloid precursor protein by
the transmembrane aspartic protease BACE".
RL Science 286:735-741(1999).
CC - FUNCTION: Responsible for the proteolytic processing of the
CC amyloid precursor protein (APP). Cleaves at the amino terminus of
CC the A-beta peptide sequence, between residues 671 and 672 of APP,
CC leads to the generation and extracellular release of beta-cleaved
CC soluble APP, and a corresponding cell-associated carboxy-terminal
CC fragment which is later released by gamma-secretase (By
CC similarity).
CC CATALYTIC ACTIVITY: Broad endopeptidase specificity. Cleaves Glu-
CC Val-Ileu-Leu-Ileu-Ala-Glu-Phe in the Swedish variant of
CC Alzheimer's amyloid precursor protein.
CC - SUBCELLULAR LOCATION: Type I membrane protein.
CC - SIMILARITY: Belongs to the peptidase A1 family.

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CC or send an email to license@sib-sib.ch).
CC -----
CC EMBL: AF190727; AAF04144.1; -.
CC HSSP: P56817; 1MAH.
DR MEROPS; A01.004; -.
DR RGD; 2191; BACE.
DR InterPro: IPR009119; Pept_A1_BACE.
DR InterPro: IPR009120; Pept_A1_BACE1.
DR InterPro: IPR001969; Pept_Asp_AS.
DR InterPro: IPR009007; Pept_AspArtic.
DR InterPro: IPR001461; Peptidase_A1.
DR Pfam; PF00026; Asp. 1.
DR PRINTS; PRO1816; BACE1.
DR PRINTS; PRO1815; BACEFAMILY.
DR PROSITE; PS00141; ASP_PROTEASE; 1.
DR PROSITE; PRO0792; PEPsin.
KW Aspartyl protease; Glycoprotein; Hydrolase; Signal; Transmembrane;
KM Zymogen.
FT SIGNAL 1 21 Potential.
FT PROPEP 22 45 Potential.
FT CHAIN 46 501 Beta-secretase 1.
FT DOMAIN 22 457 Extracellular (Potential).
FT TRANSMEM 458 478 Potential.
FT DOMAIN 479 501 Cytoplasmic (Potential).
FT ACT_SITE 93 93 By similarity.
FT ACT_SITE 289 289 By similarity.
FT DISULFID 216 420 By similarity.
FT DISULFID 278 443 By similarity.
FT DISULFID 330 380 By similarity.
FT CARBOHYD 153 153 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 172 172 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 223 223 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 354 354 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 501 AA; 55806 MW; 24B4458C8B87DE3 CRC64;

Query Match 100.0%; Score 144; DB 1; Length 501;
Best Local Similarity 100.0%; Pred. No. 2,4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTQILNIVDTGSSNPAV 28
Db 74 GYVEMTVGSPPTQILNIVDTGSSNPAV 101
|||||
RESULT 6
ID Q81YC8 PRELIMINARY; PRT; 501 AA.
AC Q81YC8;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Beta-site APP-cleaving enzyme 1, isoform A preproprotein.
GN Name=BACE1;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RX MEDLINE=22386257; Pubmed=12477932; DOI=10.1073/pnas.242603899;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altshuler S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang U., Helen F.,
RA Datchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,

RA Stopleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Uedin T.B., Toshiyuki S., Carninci P., Prange C.C.,
 RA Rana S.S., Loguettano N.A., Peters G.J., Abrahamson R.D., Mullany S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Villalón D.K., Morley K.C., Hale S., García A.M., Gay L.J., Huilyk S.W.,
 RA Fahy J., Helton E., Kettelman M., Madan A., Rodríguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakeley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butlerfield Y.S.,
 RA Krywinski M.I., Skalela U., Smallus D.E., Scherch A., Schein J.E.,
 RA Jones S.J., Maira M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences."
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RA Strauberg R.;
 RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: Belongs to peptidase family A1.
 DR EMBL; BC036084; AA036084.1; -.
 DR HSSP; P56817; 1PKN.
 DR GO; GO:0005768; C:endosome; ISS.
 DR GO; GO:0005794; C:Golgi apparatus; ISS.
 DR GO; GO:0016021; C:integral to membrane; ISS.
 DR GO; GO:0004190; F:aspartic-type endopeptidase activity; ISS.
 DR GO; GO:0050435; P:beta-amyloid metabolism; ISS.
 DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro; IPR001461; Peptidase A1.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009120; Pept_A1_BACE.
 DR InterPro; IPR009007; Pept_Aspartic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PRINTS; PRO1816; BACEFAMILY.
 DR PRINTS; PRO1815; BACEFAMILY.
 DR PRINTS; PRO0792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.
 KM Apatyrl protease; Hydrolase; Protease.
 SQ SEQUENCE 501 AA; 55823 MW; 7685955C5517EB7 CRC64;
 QY 1 GYVENTVSPPTLTILVDTGSSNFAV 28
 Db 74 GYVENTVSPPTLTILVDTGSSNFAV 101
 RESULT 7
 OBB0Y4 PRELIMINARY; PRT; 501 AA.
 ID OBB0Y4;
 AC OBB0Y4;
 DT 01-MAR-2003 (TREMBLrel. 23, Created)
 DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
 DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
 DE Mus musculus adult male corpora quadrigemina cDNA, RIKEN full-length
 DE enriched library, clone:B230346M13 product:beta-site APP cleaving
 DE enzyme, full insert sequence.
 GN Name=Bace1; Synonyms=Bace;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxId=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=99279253; PubMed=10349636; DOI=10.1016/S0076-6879(99)03004-9;
 RA Carninci P., Hayashizaki Y.;
 RT "High-efficiency full-length cDNA cloning."
 RL Meth. Enzymol. 303:19-44(1999).
 RN [2]

RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=21085660; PubMed=11217851; DOI=10.1038/35055500;
 RA RIKEN FANTOM Consortium;
 RT "Functional annotation of a full-length mouse cDNA collection."
 RN Nature 409:685-690(2001).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RA The FANTOM Consortium,
 RA the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs."
 RL Nature 420:563-573(2002).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=20499374; PubMed=11042159; DOI=10.1101/gr.145100.
 RA Carninci P., Shibata Y., Hayatsu N., Sugahara Y., Shibata K., Itoh M.,
 RA Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
 RT "Normalization and subtration of cap-trapper-selected cDNAs to
 RT prepare full-length cDNA libraries for rapid discovery of new genes."
 RL Genome Res. 10:1617-1630(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RX MEDLINE=20530913; PubMed=11076861; DOI=10.1101/gr.152600;
 RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
 RA Kono H., Akiyama Y., Nishi K., Kitamura T., Tashiro H., Itoh M.,
 RA Sumi N., Ishii Y., Nakamura S., Hazama M., Nishine T., Harada A.,
 RA Yamamoto R., Matsumoto H., Sakaguchi S., Ikegami T., Kashiwagi K.,
 RA Fujiwara S., Inoue K., Togawa Y., Izawa M., Ohara E., Watabiki M.,
 RA Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsumoto S., Kawai J.,
 RA "RIKEN integrated sequence analysis (RISA) system-384-format
 RT sequencing pipeline with 384 multicapillary sequencer."
 RL Genome Res. 10:1757-1771(2000).
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Corpora quadrigemina;
 RA Adachi J., Aizawa K., Akimura T., Arakawa T., Bono H., Carninci P.,
 RA Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
 RA Hayashida K., Hayatsu N., Hiramoto K., Hiraoka T., Hirozawa T.,
 RA Hori F., Imotani K., Ishii Y., Itoh M., Kagawa I., Kasukawa T.,
 RA Katoh H., Kawai J., Kojima Y., Kondo S., Kono H., Kouda M., Koya S.,
 RA Kurihara C., Matsuyama T., Miyazaki A., Murata M., Nakamura M.,
 RA Nishi K., Nomura K., Numazaki R., Ono M., Ohsato N., Okazaki Y.,
 RA Saito R., Saitoh H., Sakai C., Sakai K., Sakazume N., Sano H.,
 RA Sasaki D., Shibata K., Shinagawa A., Shiraki T., Sogabe Y., Tagami M.,
 RA Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
 RA Tomaru A., Toya T., Yasunishi A., Muramatsu M., Hayashizaki Y.,
 RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: Belongs to peptidase family A1.
 DR EMBL; AK046175; BAC32620.1; -.
 DR HSSP; P56817; 1PKN.
 DR MGD; MG11346542; Bace1.
 DR GO; GO:0005768; C:endosome; ISS.
 DR GO; GO:0005615; C:extracellular space; TMS.
 DR GO; GO:0005794; C:Golgi apparatus; ISS.
 DR GO; GO:0016021; C:integral to membrane; ISS.
 DR GO; GO:0004190; F:aspartic-type endopeptidase activity; ISS.
 DR GO; GO:0050435; P:beta-amyloid metabolism; ISS.
 DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR InterPro; IPR001461; Peptidase A1.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009120; Pept_A1_BACE.
 DR InterPro; IPR009007; Pept_Aspartic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PRINTS; PRO1816; BACE1.
 DR PRINTS; PRO1815; BACEFAMILY.
 DR PRINTS; PRO0792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 1.
 KM Apatyrl protease; Hydrolase; Protease.

SEQ SEQUENCE 501 AA; 55816 MW; C085513145E024E CRC64;
Query Match 100.0%; Score 144; DB 2; Length 501;
Best Local Similarity 100.0%; Pred. No. 2, 4e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
Db 74 GYVEMTVGSPPTLNIIVDTGSSNFAV 101
RESULT 8
Q9ULS1 PRELIMINARY; PRT; 532 AA.
ID Q9ULS1
AC Q9ULS1
DT 01-MAR-2000 (TrEMBLrel. 13, Created)
DT 01-OCT-2001 (TrEMBLrel. 18, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE KIAA1149 protein (Fragment).
GN Name=KIAA1149;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain; N.A.
RX MEDLINE=20039618; PubMed=10574461;
RA Hirosewa M., Nagase T., Ishikawa K., Kikuno R., Nomura N., Ohara O.;
RT "Characterization of cDNA clones selected by the Genemark analysis
RT from size-fractionated cDNA libraries from human brain.";
RL DNA Res. 6:329-336(1999).
CC -1- SIMILARITY: Belongs to peptidase family A1.
DR EMBL; AB032975; BAA8463.2; -;
DR HSSP; P56817; 1PKN.
DR GO; GO:0005768; C:Endosome; ISS.
DR GO; GO:0005794; C:Golgi apparatus; ISS.
DR GO; GO:0016021; C:Integral to membrane; ISS.
DR GO; GO:0004190; F:Aspartic-type endopeptidase activity; ISS.
DR GO; GO:0005035; P:beta-amyloid metabolism; ISS.
DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
DR InterPro: IPR001461; Peptidase A1.
DR InterPro: IPR009119; Pept. A1_BACE.
DR InterPro: IPR009120; Pept. A1_BACE.
DR InterPro: IPR009007; Pept. Aspartic.
DR InterPro: IPR001969; Pept. Asp_AS.
DR PRINTS; PR01816; BACE1.
DR PRINTS; PR01815; BACEFAMILY.
DR PRINTS; PR00792; PEPsin.
DR PROSITE; PS00141; ASP_PROTEASE; 1.
KW Aspartyl protease; Hydrolyase; Protease.
FT NON_TER 1
SQ SEQUENCE 532 AA; 58720 MW; 98B135D0D5FBD2E8 CRC64;
Query Match 100.0%; Score 144; DB 2; Length 532;
Best Local Similarity 100.0%; Pred. No. 2, 6e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28
Db 105 GYVEMTVGSPPTLNIIVDTGSSNFAV 132
RESULT 9
Q8C7R1 PRELIMINARY; PRT; 501 AA.
ID Q8C7R1
AC Q8C7R1
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Mus musculus 12 days embryo spinal cord cDNA, RIKEN full-length
DE enriched library, clone: C530008K17 product: beta-site App cleaving
DE enzyme, full insert sequence.

GN Name=Bacel; Synonyms=Bace;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Spinal cord;
RX MEDLINE=99279253; PubMed=10349636; DOI=10.1016/S0076-6879(99)03004-9;
RA Carninci P., Hayashizaki Y.;
RT "High-efficiency full-length cDNA cloning.";
RL Mech. Enzymol. 303:19-44(1999).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Spinal cord;
RX MEDLINE=21085660; PubMed=11217851; DOI=10.1038/35055500;
RA RIKEN FANTOM Consortium;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Spinal cord;
RA The FANTOM Consortium;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573(2002).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Spinal cord;
RX MEDLINE=20499374; PubMed=11042159; DOI=10.1101/97.145100;
RA Carninci P., Shibata Y., Hayatsu N., Sugahara Y., Shibata K., Itoh M.,
RA Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
RT "Normalization and subtractions of cap-trapper-selected cDNAs to
RT prepare full-length cDNA libraries for rapid discovery of new genes.";
RL Genome Res. 10:1617-1630(2000).
RN [5]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Spinal cord;
RX MEDLINE=20530913; PubMed=11076861; DOI=10.1101/gr.152600;
RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
RA Kono H., Akiyama J., Nishi K., Katsunai T., Tashiro H., Itoh M.,
RA Sumi N., Ishii Y., Nakamura S., Hazama M., Nishine T., Harada A.,
RA Yamamoto R., Matsunoto H., Sakaguchi S., Ikegami T., Kashiwagi K.,
RA Fujiwaki S., Inoue K., Togawa Y., Izawa Y., Ohara E., Wataniki M.,
RA Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsunaga S., Kawai J.,
RA Okazaki Y., Muramatsu M., Inoue Y., Kira A., Hayashizaki Y.;
RT "RIKEN integrated sequence analysis (RISA) system-384-format
RT sequencing pipeline with 384 multicapillary sequencer.";
RL Genome Res. 10:1757-1771(2000).
RN [6]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Spinal cord;
RA Adachi J., Aizawa K., Akimura T., Arakawa T., Bono H., Carninci P.,
RA Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
RA Hayashida K., Hayatsu N., Hiramoto K., Hiraoka T., Hirozane T.,
RA Hori F., Imclani K., Ishii Y., Itoh M., Kigawa I., Kasukawa T.,
RA Katon H., Kawai J., Kojima Y., Kondo S., Kono H., Kouda M., Koya S.,
RA Kurihara C., Matsuyama T., Miyazaki A., Murata M., Nakamura M.,
RA Nishi K., Nomura K., Numazaki R., Ohno M., Ono N., Okazaki Y.,
RA Saito R., Saitoh H., Sakai C., Sakai K., Sakazume N., Sano H.,
RA Sasaki D., Shibata K., Shinagawa A., Shiraki T., Sogabe Y., Tagami M.,
RA Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
RA Tomaru A., Toyota T., Yasunishi A., Muramatsu M., Hayashizaki Y.;
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: Belongs to peptidase family A1.
DR EMBL; AK049626; BAC33844.1; -;
DR HSSP; P56817; 1PKN.
DR MGD; MGI:1346542; Bacel.
DR GO; GO:0005768; C:Endosome; ISS.
DR GO; GO:0005615; C:extracellular space; TAS.
DR GO; GO:0005794; C:Golgi apparatus; ISS.
DR GO; GO:0016021; C:Integral to membrane; ISS.

Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
 RL EMBL: BC055989; AAH55989.1; -
 DR HSSP; P56817; 1FKN.
 DR GO; GO:0004194; F:pepsin A activity; IEA.
 DR GO; GO:0008233; F:peptidase activity; IEA.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro; IPR009007; Pept_AspArtic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PROSITE; PS00141; ASP_PROTEASE; 2.
 DR SEQUENCE 500 AA; 54722 MW; 10FL6756CAFDDC0B CRC64;
 Query Match 84.7%; Score 122; DB 2; Length 500;
 Best Local Similarity 78.6%; Pred. No. 66-10;
 Matches 22; Conservative 4; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 GYVENTVGSPPQTNIIVDTGSSNFAV 28
 Db 75 GYVELLIGSPQKNVILVDTGSSNFAV 102
 RESULT 12
 Q6PB20 PRELIMINARY; PRT; 499 AA.
 AC Q6PB20;
 DT 05-JUL-2004 (TrEMBLrel. 27, Created)
 DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
 DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
 DE MGC68482 protein.
 GN Name=MGC68482;
 OS Xenopus laevis (African clawed frog).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipidoidea; Pipidae;
 OC Xenopodinae; Xenopus.
 OX NCBI_TaxID=83355;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Kidney;
 RX MEDLINE=22388557; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altshul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldi M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Ustun T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abirmason R.D., Mullany S.J.,
 RA Bosak S.A., McGowan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Morley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahy J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
 RA Blakeley R.W., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
 RA Krzywinski M.I., Skalka U., Smallos D.E., Schnerch A., Schein J.E.,
 RA Jones S.J., Marra M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences."
 RT Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Kidney;
 RX MEDLINE=22341132; PubMed=12454917; DOI=10.1002/dvdy.10174;
 RA Klein S.L., Strausberg R.L., Wagner L., Pontius J., Clifton S.W.,
 RA Richardson P.,
 RT "Genetic and genomic tools for Xenopus research: The NIH Xenopus
 RT initiative."
 RT Dev. Dyn. 225:384-391(2002).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Kidney;
 RX Klein S., Strausberg R.,
 RT Submitted (OCT-2003) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: Belongs to peptidase family A1.

DR EMBL: BC059963; AAH59963.1; -
 DR HSSP; P20142; 1AVF.
 DR GO; GO:0009049; F:aspartic-type signal peptidase activity; IEA.
 DR GO; GO:0004194; F:pepsin A activity; IEA.
 DR GO; GO:0008233; F:peptidase activity; IEA.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro; IPR001461; Peptidase_A1.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009121; Pept_A1_BACE2.
 DR InterPro; IPR009007; Pept_AspArtic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PRINTS; PR01817; BACE2.
 DR PRINTS; PR01815; BACEFAMILY.
 DR PRINTS; PR00792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 2.
 DR Aspartyl protease; Hydrolase; Protease.
 DR SEQUENCE 499 AA; 54803 MW; E846674A5DF68AF CRC64;
 Query Match 83.3%; Score 120; DB 2; Length 499;
 Best Local Similarity 75.0%; Pred. No. 1,2e-09;
 Matches 21; Conservative 5; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 GYVENTVGSPPQTNIIVDTGSSNFAV 28
 Db 74 GYVELLIGTTPQKNVILVDTGSSNFAV 101
 RESULT 13
 Q9NZL1 PRELIMINARY; PRT; 396 AA.
 AC Q9NZL1;
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Aspartyl protease.
 GN Name=BACE2;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20422477; PubMed=10965118;
 RA Solans A., Estivill X., de la Luna S.,
 RT "A new aspartyl protease on 21q22.3, BACE2, is highly similar to
 RT Alzheimer's amyloid precursor protein beta-secretase."
 RT Cytogenet. Cell Genet. 89:177-184(2000).
 CC -1- SIMILARITY: Belongs to peptidase family A1.
 DR EMBL: AF188277; AAF35836.1; -
 DR HSSP; P56817; 1FKN.
 DR GO; GO:0016021; C:integral to membrane; ISS.
 DR GO; GO:0006509; P:membrane protein ectodomain proteolysis; ISS.
 DR GO; GO:0042985; P:negative regulation of amyloid precursor pr. . .; ISS.
 DR GO; GO:0016486; P:peptide hormone processing; ISS.
 DR InterPro; IPR001461; Peptidase_A1.
 DR InterPro; IPR009119; Pept_A1_BACE.
 DR InterPro; IPR009121; Pept_A1_BACE2.
 DR InterPro; IPR009007; Pept_AspArtic.
 DR InterPro; IPR001969; Pept_Asp_AS.
 DR PRINTS; PR01815; BACE2.
 DR PRINTS; PR00792; PEPsin.
 DR PROSITE; PS00141; ASP_PROTEASE; 2.
 DR Aspartyl protease; Hydrolase; Protease.
 DR SEQUENCE 396 AA; 43013 MW; 5023A7AF391CEAC9 CRC64;
 Query Match 82.6%; Score 119; DB 2; Length 396;
 Best Local Similarity 78.6%; Pred. No. 1,3e-09;
 Matches 22; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
 Qy 1 GYVENTVGSPPQTNIIVDTGSSNFAV 28
 Db 91 GYVELLIGTTPQKNVILVDTGSSNFAV 118

CC therapy. (1) can be used for producing preparations of homogenously
CC processed BACE that may be used for e.g. studying or treating diseases
CC such as Alzheimer's disease or Down's syndrome. The human BACE1 gene is
CC located on chromosome 11, more specifically to 11q23.2-23.3. The present
CC sequence represents a human BACE1 isoform A protease domain amino acid
CC sequence, which is used in the exemplification of the present invention.
XX
SQ Sequence 28 AA;

Query Match 100.0%; Score 144; DB 8; Length 28;
Best Local Similarity 100.0%; Pred. No. 6,9e-15;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTLTNLIVDSSNFAV 28
Db 1 GYVEMTVGSPPTLTNLIVDSSNFAV 28
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1 GYVEMTVGSPPTLTNLIVDSSNFAV 28

RESULT 2
AAU23068
ID AAU23068 standard; protein; 387 AA.
XX
AC AAU23068;
XX
DT 17-DEC-2001 (first entry)
XX
DE Novel human enzyme polypeptide #154.
XX
KW Human; oxidoreductase enzyme; transferase; hydrolase; lyase; isomerase;
KW ligase; hyperproliferative disorder; immunodeficiency disorder;
KW autoimmune disorder; neurological disorder; metabolic disorder;
KW inflammatory disorder; cardiovascular disorder; reproductive disorder;
KW blood-related disorder; infectious disorder; cytostatic; anti arthritic;
KW nephrotropic; anticoagulant.
XX
OS Homo sapiens.
XX
PN MO200155301-A2.
XX
PD 02-AUG-2001.
XX
PF 17-JAN-2001; 2001WO-US001239.
XX
PR 31-JAN-2000; 2000US-0179065P.
PR 04-FEB-2000; 2000US-0180628P.
PR 24-FEB-2000; 2000US-0184664P.
PR 02-MAR-2000; 2000US-0186350P.
PR 16-MAR-2000; 2000US-0189874P.
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PR 14-AUG-2000; 2000US-0225759P.

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PR 13-OCT-2000; 2000US-0237040P.
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PR 08-NOV-2000; 2000US-0246527P.
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PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.

PR 08-NOV-2000; 2000US-0246613P.
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PR 17-NOV-2000; 2000US-0249209P.
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PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
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PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
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PR 17-NOV-2000; 2000US-0249264P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
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PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
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PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
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PR 08-DEC-2000; 2000US-0251989P.
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PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Rosen CA, Barash SC, Ruben SM;
XX
DR WPI; 2001-465566/50.
XX N-PSDB; AAS40938.
XX
PT Novel polypeptides and polynucleotides useful for diagnosing, preventing,
XX treating neural, immune system, muscular, reproductive, pulmonary,
XX cardiovascular, renal, proliferative disorders and cancerous diseases.
XX
PS Claim 11; SEQ ID NO 1064; 1180bp; English.
XX
XX The present invention relates to the isolation of novel human enzyme
XX polypeptides, and the cDNA (AAS40795-AAS41684) and genomic sequences
XX encoding them. The enzyme polypeptides of the invention may comprise the
XX functional classes of oxidoreductases, transferases, hydrolases, lyases,
XX isomerases or ligases. The sequences of the invention are useful in the
XX diagnosis, treatment, prevention and/or prognosis of a wide range of
XX disorders including hyperproliferative disorders (e.g. cancer),
XX immunodeficiency disorders (e.g. AIDS) autoimmune disorders (e.g.
XX arthritis), neurological disorders (e.g. Alzheimer's disease), metabolic
XX disorders (e.g. phenylketonuria), inflammatory disorders (e.g. asthma),
XX cardiovascular disorders (e.g. atherosclerosis), blood-related disorders
XX (e.g. haemophilia), reproductive disorders (e.g. infertility) and
XX infectious disorders (e.g. influenza). The polynucleotides of the
XX invention can also be used in gene therapy. AAU22915-AAU23814 represent
XX the novel human enzyme polypeptides of the invention. Note: The sequence
XX data for this patent did not form part of the printed specification, but
XX was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 387 AA;
XX
Query Match 100.0%; Score 144; DB 4; Length 387;
Best Local Similarity 100.0%; Pred. No. 1.6e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GYVEMTVGSPPTLNIIVDTGSSNFAV 28

Db |||||
10 GYVEMTVGSPPTLNIIVDTGSSNFAV 37
RESULT 3
AAU23069
ID AAU23069 standard; protein; 390 AA.
XX
XX AAU23069;
AC
XX
DT 17-DEC-2001 (first entry)
XX
DE Novel human enzyme polypeptide #155.
XX
XX Human; oxidoreductase enzyme; transferase; hydrolase; lyase; isomerase;
XX ligase; hyperproliferative disorder; immunodeficiency disorder;
XX autoimmune disorder; neurological disorder; metabolic disorder;
XX inflammatory disorder; cardiovascular disorder; reproductive disorder;
XX blood-related disorder; infectious disorder; cytostatic; anti arthritic;
XX nephrotropic; anticoagulant.
XX
XX Homo sapiens.
XX
XX WO200155301-A2.
XX
XX 02-AUG-2001.
XX
XX 17-JUN-2001; 2001WO-US001239.
XX
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184648P.
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XX 06-SEP-2000; 2000US-0230438P.

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PR 17-NOV-2000; 2000US-0249245P.
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 PR 05-DEC-2000; 2000US-0251988P.
 PR 05-DEC-2000; 2000US-0251989P.
 PR 06-DEC-2000; 2000US-0251479P.
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 PR 08-DEC-2000; 2000US-0251990P.
 PR 11-DEC-2000; 2000US-0254097P.
 PR 05-JAN-2001; 2001US-0259678P.
 PA (HUMA-) HUMAN GENOME SCI INC.
 PI Rosen CA, Barash SC, Ruben SM;
 XX WPI, 2001-465566/50.
 DR N-PSDB; AAS40939.
 XX
 PT Novel polypeptides and polynucleotides useful for diagnosing, preventing,
 PT treating neutral, immune system, muscular, reproductive, pulmonary,
 PT cardiovascular, renal, proliferative disorders and cancerous diseases.
 PS
 XX Claim 11; SEQ ID NO 1065; 1180bp; English.
 CC The present invention relates to the isolation of novel human enzyme
 CC polypeptides, and the cDNA (AAS40785-AAS41684) and genomic sequences
 CC encoding them. The enzyme polypeptides of the invention may comprise the
 CC functional classes of oxidoreductases, transferases, hydrolases, lyases,
 CC isomerases or ligases. The sequences of the invention are useful in the
 CC diagnosis, treatment, prevention and/or prognosis of a wide range of
 CC disorders including hyperproliferative disorders (e.g. cancer),
 CC immunodeficiency disorders (e.g. AIDS) autoimmune disorders (e.g.
 CC arthritis), neurological disorders (e.g. Alzheimer's disease), metabolic
 CC disorders (e.g. phenylketonuria), inflammatory disorders (e.g. asthma),
 CC cardiovascular disorders (e.g. atherosclerosis), blood-related disorders
 CC (e.g. haemophilia), reproductive disorders (e.g. infertility) and
 CC infectious disorders (e.g. influenza). The polynucleotides of the
 CC invention can also be used in gene therapy. AAU22915-AAU23814 represent
 CC the novel human enzyme polypeptides of the invention. Note: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 390 AA;
 Qy Query Match 100.0%; Score 144; DB 4; Length 390;
 Db Best Local Similarity 100.0%; Pred. No. 1,6e-13;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 GYVEMTVGSPPTLITLVDTGSSNFAV 28
 |||
 Db 13 GYVEMTVGSPPTLITLVDTGSSNFAV 40
 RESULT 4
 ID ADC81581 standard; protein; 391 AA.
 AC ADC81581;
 XX
 XX
 DT 01-JAN-2004 (first entry)
 XX
 XX Beta-secretase (1fkn) amino acid sequence SEQ ID NO:4.
 DE
 XX

KM	human; BACE; modification; Pro33lys; pro-enzyme.
XX	Unidentified.
OS	
XX	WO2003072733-A2.
PN	
XX	04-SEP-2003.
PD	
XX	
XX	21-FEB-2003; 2003WO-US005508.
PF	
XX	21-FEB-2002; 2002US-0358651P.
PR	
XX	(PHAA) PHARMACIA & UPJOHN CO.
PA	
XX	
PI	Chou K, Howe JW;
XX	WPI; 2003-712719/67.
DR	
XX	
PT	BACE polypeptides having Pro33lys modification, useful in determining
PT	possible mutations, which will inhibit enzyme activity, and in
XX	determining potential active site for target molecules.
XX	
PS	Disclosure; Fig 3; 38pp; English.
XX	
CC	The present invention describes an isolated polypeptide (I) comprising or
CC	consisting of a fully defined sequence of 432 amino acids (see ADG81561),
CC	and comprising human BACE having the modification Pro33lys. Also
CC	described: (1) a composition comprising an active human BACE enzyme
CC	comprising the pro-enzyme sequence of BACE having the modification
CC	Pro33lys; (2) an isolated polynucleotide comprising a sequence encoding
CC	(1); (3) an isolated polynucleotide consisting or comprising of
CC	nucleotides 70-1165 of a 1355-bp sequence (see ADG81562); (4) an
CC	expression vector comprising the polynucleotide of (2), or a
CC	polynucleotide sequence encoding a Pro33lys-BACE polypeptide, where the
CC	expression vector can produce the Pro33lys-BACE polypeptide when present
CC	in a competent host cell, when cultured under conditions that allow
CC	production; (5) a recombinant host cell comprising the expression vector;
CC	and (6) producing a (active) Pro33lys-BACE polypeptide. The BACE
CC	polypeptide having Pro33lys modification may be used in determining
CC	possible mutations, which will inhibit enzyme activity, and in
CC	determining potential active site for target molecules. The vector
CC	comprising the BACE polynucleotide is useful for producing recombinant
CC	BACE polypeptides having Pro33lys modification. The present sequence
CC	represents a beta-secretase amino acid sequence, which is used in the
CC	exemplification of the present invention.
XX	
SO	Sequence 391 AA;
Query Match	100.0%; Score 144; DB 7; Length 391;
Best Local Similarity	100.0%; Pred. No. 1.6e-13;
Matches	28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 GYVEMTWGSPPTQTLNIIIVDTGSSNFAV 28
DB	19 GYVEMTWGSPPTQTLNIIIVDTGSSNFAV 46
AD164643	standard; protein; 403 AA.
AD164643	
AD164643	
AD164643	
22-APR-2004	(first entry)
Mature human beta-secretase (BACE) protein seq id 4.	
crystal; glycosylated human beta-secretase; BACE; human beta-secretase;	
protein co-ordinate data.	
Homo sapiens.	
US2004014194-A1.	

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XX 22-JAN-2004.
PD
XX
XX 26-MAR-2003; 2003US-00400273.
PF
XX
XX 27-MAR-2002; 2002US-0367937P.
PR
XX
XX (SCHE ) SCHERING CORP.
PA
XX
XX Beyer BM, Hammond GS, Reichert P, Strickland C, Wang W, Weber PC;
PI Wong GT, Zhang L;
XX
XX WPI; 2004-167920/16.
DR
XX
XX New crystal comprising a glycosylated, human beta-secretase polypeptide,
PT useful for determining the three-dimensional structure of beta-secretase
and other related proteins.
XX
XX Claim 5; SEQ ID NO 4; 107pp; English.
PS
XX
XX The invention describes a crystal comprising a glycosylated, human beta-
CC secretase polypeptide characterised by structural coordinates comprising
CC a root mean square deviation of conserved residue backbone atoms of less
CC than 1.5 Angstrom when superimposed on backbone atoms described by
CC structural coordinates. The crystal is useful for determining the three-
CC dimensional structure of beta-secretase and other related proteins. This
CC is the amino acid sequence of a mature human beta-secretase (BACE)
CC protein.
CC
SQ Sequence 403 AA;

Query Match 100.0%; Score 144; DB 8; Length 403;
Best Local Similarity 100.0%; Pred. No. 1.7e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTNLIVDGSNFAV 28
   |||||
DB 27 GYVEMTVGSPPTNLIVDGSNFAV 54

RESULT 6
AD164644
ID AD164644 standard; protein; 408 AA.
XX
XX AD164644;
AC
XX
DT 22-APR-2004 (first entry)
XX
XX Mature human beta-secretase (BACE) protein seq id 5.
XX
XX crystal; glycosylated human beta-secretase; BACE; human beta-secretase;
KW protein co-ordinate data.
XX
XX Homo sapiens.
OS
XX US2004014194-A1.
PN
XX
PD 22-JAN-2004.
XX
PF 26-MAR-2003; 2003US-00400273.
XX
XX 27-MAR-2002; 2002US-0367937P.
PR
XX
XX (SCHE ) SCHERING CORP.
PA
XX
XX Beyer BM, Hammond GS, Reichert P, Strickland C, Wang W, Weber PC;
PI Wong GT, Zhang L;
XX
XX WPI; 2004-167920/16.
DR
XX
XX New crystal comprising a glycosylated, human beta-secretase polypeptide,
PT useful for determining the three-dimensional structure of beta-secretase
and other related proteins.

```


OS Synthetic.
XX
XX WO2004011641-A2.
XX
XX 05-FEB-2004.
XX
XX 25-JUL-2003; 2003WO-GB003200.
XX
XX 26-JUL-2002; 2002US-0398681P.
XX
XX (ASTE-) ASTEX TECHNOLOGY LTD.
XX
XX Vuillard LM, Patel SJ, Yon JR, Cleasby A, Hamilton BJ, Shah A;
XX WPI; 2004-169242/16.
XX
XX New beta site APP cleaving enzyme (BACE) protein, useful for treating or
XX preventing Alzheimer's disease or Alzheimer's-type pathology of Down's
XX syndrome.
XX
XX Claim 10; SEQ ID NO 20; 145pp; English.
XX
XX The present invention relates to a beta site APP cleaving enzyme (BACE)
XX protein. The compound or the composition is useful in medicine and the
XX BACE crystal structure is useful for drug discovery. The BACE protein,
XX compounds, pharmaceutical compositions, medicament, drug or other
XX composition comprising the compound is useful for treating or preventing
XX Alzheimer's disease or Alzheimer's-type pathology of Down's syndrome. The
XX present sequence represents the DNA sequence for a BACE protein.
XX
XX Sequence 411 AA;
XX
XX

Query Match 100.0%; Score 144; DB 8; Length 411;
Best Local Similarity 100.0%; Pred. No. 1.7e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GYVENTVGSPPQTLNIIIVDTGSSNPAV 28
DB 32 GYVENTVGSPPQTLNIIIVDTGSSNPAV 59

RESULT 10
ABR61930
ID ABR61930 standard; protein; 414 AA.
XX
XX ABR61930;
XX
XX 12-SEP-2003 (first entry)
XX
XX Human promemapsin 2 protein fragment.
XX
XX Memapsin 1; neurotrophic; neuroprotective; memapsin 2; beta secretase;
XX beta-amyloid protein; Alzheimer's disease; promemapsin 2; human.
XX
XX Homo sapiens.
XX
XX WO2003039454-A2.
XX
XX 15-MAY-2003.
XX
XX 23-OCT-2002; 2002WO-US034324.
XX
XX 23-OCT-2001; 2001US-0335952P.
XX 27-NOV-2001; 2001US-033545P.
XX 14-JAN-2002; 2002US-0348464P.
XX 20-JUN-2002; 2002US-0390804P.
XX 19-JUL-2002; 2002US-0397557P.
XX 19-JUL-2002; 2002US-0397619P.
XX
XX (OKLA-) OKLAHOMA MEDICAL RES FOUND.
XX (UNIT) UNIV ILLINOIS FOUND.
XX

PI Ghosh AK, Tang J, Bilcer G, Chang W, Hong L, Koelsch G, Loy J;
PI Turner RT;
XX
XX WPI; 2003-541410/51.
XX
XX New peptide compounds are memapsin beta secretase inhibitors used for
XX treating Alzheimer's disease.
XX
XX Example; Fig 12; 407pp; English.
XX
XX The invention relates to peptide compounds of specified formula. The
XX compounds exhibit memapsin 2-beta secretase inhibitory activity relative
XX to memapsin 1-beta secretase and reduce the accumulation of beta-amyloid
XX protein. The compounds can be used for treating Alzheimer's disease. The
XX present sequence represents a human promemapsin 2 protein fragment used
XX in crystal structures
XX
XX Sequence 414 AA;
XX
XX

Query Match 100.0%; Score 144; DB 6; Length 414;
Best Local Similarity 100.0%; Pred. No. 1.7e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GYVENTVGSPPQTLNIIIVDTGSSNPAV 28
DB 34 GYVENTVGSPPQTLNIIIVDTGSSNPAV 61

RESULT 11
ADC72735
ID ADC72735 standard; protein; 414 AA.
XX
XX ADC72735;
XX
XX 18-DEC-2003 (first entry)
XX
XX Human beta-site aspartyl protease cleaving enzyme catalytic domain.
XX
XX neuroprotective; neurotrophic; crystalline; Beta-site APP cleaving enzyme;
XX BACE; aspartyl protease; Alzheimer's disease; protein co-ordinate data.
XX
XX Homo sapiens.
XX
XX WO2003012089-A2.
XX
XX 13-FEB-2003.
XX
XX 26-JUL-2002; 2002WO-GB003461.
XX
XX 26-JUL-2001; 2001US-0308366P.
XX
XX (ASTE-) ASTEX TECHNOLOGY LTD.
XX (JANC) JANSSEN PHARM NV.
XX
XX Yon J, Cleasby A, Bruinzeel WD, Masure SLJ, Tickle I, Sharff A;
XX WPI; 2003-239524/23.
XX N-PSDB; ADC72736.
XX
XX New Beta-site APP cleaving enzyme (BACE) proteins and protein crystal,
XX useful in designing compounds that inhibit or modulate BACE, in drug
XX screening assays, and in identifying receptors.
XX
XX Disclosure; Fig 2a; 272pp; English.
XX
XX The invention relates to a new crystalline form of Beta-site APP cleaving
XX enzyme (BACE) or its functional portion having an active site containing
XX one or more ligands other than the natural substrate or the substrate
XX that occurs naturally or physiologically within the active site.
XX Inhibitors of BACE protein or its functional portion is useful for
XX preparing a composition or medicament for inhibiting BACE or the
XX production of A-beta or its fragments, and in therapy for treating
XX Alzheimer's disease. The BACE crystals and proteins may be used to design
XX

CC compounds that inhibit or modulate BACE, in drug screening assays, and in
CC identifying receptors. This sequence represents a fragment of the full
CC length BACE protein from amino acid 76 to the C-terminus.
XX
SQ Sequence 414 AA;

Query Match 100.0%; Score 144; DB 7; Length 414;
Best Local Similarity 100.0%; Pred. No. 1.7e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTQNLILVDTGSSNPAV 28
DB 29 GYVEMTVGSPPTQNLILVDTGSSNPAV 56

RESULT 12
AAB07899
ID AAB07899 standard; protein; 415 AA.

XX AAB07899;

XX 14-NOV-2000 (first entry)

DE Amino acid sequence of a human beta-secretase enzyme fragment.

XX Beta-secretase; beta-amyloid precursor protein; beta-amyloid peptide;
KW amyloid plaque component; Alzheimer's disease; amyloidogenic disease;
KW inhibitor.

XX Homo sapiens.

XX WO200047618-A2.

XX 17-AUG-2000.

XX 10-FEB-2000; 2000WO-US003819.

XX 10-FEB-1999; 99US-0119571P.

XX 15-JUN-1999; 99US-0139172P.

XX (ELAN-) ELAN PHARM INC.

PI Anderson JP, Basi G, Doane MT, Frigon N, John V, Power M;
PI Simha S, Tatsuno G, Tung J, Wang S, Mcconlogue L;

XX WPI; 2000-533011/48.

PT Purified beta-secretase protein used in assays to discover inhibitors
PT which can be used for the treatment of amyloidogenic diseases e.g.
XX Alzheimer's disease.

XX Claim 10; Fig 3B; 121pp; English.

CC The specification describes a beta-secretase enzyme. The enzyme cleaves
CC beta-amyloid precursor protein to produce beta-amyloid peptide. This
CC enzyme is therefore implicated in the production of amyloid plaque
CC components which accumulate in the brains of individuals afflicted with
CC Alzheimer's disease. Inhibitors of beta-secretase are administered to a
CC mammalian subject e.g. with Alzheimer's disease or Alzheimer's disease-
CC like pathology to test if they maintain or improve cognitive ability or
CC reduce the plaque burden. The compounds are used for the treatment of
CC amyloidogenic diseases e.g. Alzheimer's disease. The present sequence
CC represents a human beta-secretase enzyme fragment

XX Sequence 415 AA;

Query Match 100.0%; Score 144; DB 3; Length 415;
Best Local Similarity 100.0%; Pred. No. 1.8e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTQNLILVDTGSSNPAV 28
DB 29 GYVEMTVGSPPTQNLILVDTGSSNPAV 56

RESULT 13
ADJ57792
ID ADJ57792 standard; protein; 417 AA.

XX ADJ57792;

XX 06-MAY-2004 (first entry)

DE BACE N-Q R56KR57 crystallised protein.

XX beta site APP cleaving enzyme; BACE; Nootropic; Neuroprotective;
KW Alzheimer's disease.

XX Synthetic.

XX WO2004011641-A2.

XX 05-FEB-2004.

XX 25-JUL-2003; 2003WO-GB003200.

XX 26-JUL-2002; 2002US-0398681P.

XX (ASTE-) ASTEX TECHNOLOGY LTD.

PI Vulliamd LMM, Patel SJ, Yon JR, Cleasby A, Hamilton BJ, Shah A;

XX WPI; 2004-169242/16.

PT New beta site APP cleaving enzyme (BACE) protein, useful for treating or
PT preventing Alzheimer's disease or Alzheimer's-type pathology of Down's
PT syndrome.

XX Claim 10; SEQ ID NO 21; 145pp; English.

CC The present invention relates to a beta site APP cleaving enzyme (BACE)
CC protein. The compound or the composition is useful in medicine and the
CC BACE crystal structure is useful for drug discovery. The BACE protein,
CC compounds, pharmaceutical compositions, medicament, drug or other
CC composition comprising the compound is useful for treating or preventing
CC Alzheimer's disease or Alzheimer's-type pathology of Down's syndrome. The
CC present sequence represents the DNA sequence for a BACE protein.

XX Sequence 417 AA;

Query Match 100.0%; Score 144; DB 8; Length 417;
Best Local Similarity 100.0%; Pred. No. 1.8e-13;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GYVEMTVGSPPTQNLILVDTGSSNPAV 28
DB 32 GYVEMTVGSPPTQNLILVDTGSSNPAV 59

RESULT 14
AAH8437
ID AAH8437 standard; protein; 425 AA.

XX AAH8437;

XX 03-AUG-2000 (first entry)

DE Human Asp2 amino acid sequence containing proteolytic cleavage site.

XX Aspartyl protease; aspartase; amyloid precursor protein; APP; Asp 2;

XX Alzheimer's disease; beta secretase site.

XX Homo sapiens.

XX WO200017369-A2.

PD 30-MAR-2000.
 XX
 PF 23-SEP-1999; 99WO-US020861.
 XX
 PR 24-SEP-1998; 98US-0101594P.
 XX
 PA (PHAA) PHARMACIA & UPJOHN CO.
 XX
 PI Gurney ME, Bienkowski MJ, Heinrichson RL, Parodi LA, Yan R;
 DR MPI; 2000-303209/26.
 DR N-PSDB; AAA15677.
 PT New enzyme designated human aspartase useful in research into Alzheimer's
 PT disease is capable of cleaving amyloid protein precursor at the beta
 PT secretase site to produce amyloid beta peptide.
 XX
 PS Example 9; Page 166-168; 183pp; English.
 XX
 CC This sequence represents a modified version of the human aspartase 2
 CC (Asp2) nucleotide sequence. The sequence is used in the bacterial
 CC expression of human Asp2L. The invention relates to a protease (e.g.
 CC Asp2) capable of cleaving the beta secretase site of amyloid precursor
 CC protein (APP). The protease contains a sequence encoding the amino acid
 CC sequence DTG and a sequence encoding DSG or DTG separated by 100-300
 CC amino acids. When mutated the APP gene causes an autosomal dominant form
 CC of Alzheimer's disease. APP localises to the cell surface membrane and
 CC have a single C-terminal transmembrane domain. Proteolytic processing of
 CC APP produces the amyloid beta protein, which is possibly very important
 CC in Alzheimer's disease. The invention includes a nucleotide sequence
 CC encoding the protease, a vector containing the nucleotide sequence, and a
 CC cell line comprising the vector. Methods for screening for inhibitors of
 CC beta secretase activity are also given in the invention. The human
 CC aspartase protein and nucleotide sequences and the methods for
 CC identifying inhibitors of the protease, are useful in the treatment of
 CC and research in to Alzheimer's disease
 XX
 SQ Sequence 425 AA;
 Query Match 100.0%; Score 144; DB 3; Length 425;
 Best Local Similarity 100.0%; Pred. No. 1.8e-13;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GYVEMTVGSPPTQTLNLTVDGSSNFAV 28
 DB 46 GYVEMTVGSPPTQTLNLTVDGSSNFAV 73
 RESULT 15
 AAU07214
 ID AAU07214 standard; protein; 425 AA.
 XX
 AC AAU07214;
 XX
 DT 11-SEP-2003 (revised)
 DT 24-OCT-2001 (first entry)
 XX
 DE T7-caspase-caspase 8-human aspartyl protease 2a deltaTM.
 XX
 KM Human; aspartyl protease 1; Asp-1; neurotrophic; neuroprotective;
 KM aspartyl protease 2; Asp2; amyloid protein precursor; APP;
 KM beta-secretase; Alzheimer's disease.
 XX
 OS Homo sapiens.
 OS Enterobacteria phage T7.
 XX
 PN WO200149097-A2.
 XX
 PD 12-JUL-2001.
 XX
 PF 09-MAY-2001; 2001WO-IB000797.
 XX
 PR 09-MAY-2001; 2001WO-IB000797.

XX
 PA (BIEN/) BIENKOWSKI M J.
 PA (GURN/) GURNEY M E.
 PA (HEIN/) HEINRICHSON R L.
 PA (PARO/) PARODI L A.
 PA (YANR/) YAN R.
 XX
 PI Bienkowski MJ, Gurney ME, Heinrichson RL, Parodi LA, Yan R;
 DR MPI; 2001-502548/55.
 DR N-PSDB; AAS11714.
 PT Novel purified polypeptide comprising fragment of mammalian aspartyl
 PT protease 2, lacking Asp2 transmembrane domain and retaining beta
 PT secretase activity of Asp2 useful for identifying inhibitors of Asp2
 PT activity.
 XX
 PS Example 9; Page 158-159; 185pp; English.
 XX
 CC The invention relates to a novel purified polypeptide comprising a
 CC fragment of mammalian aspartyl protease 2 (Asp2) protein which lacks the
 CC Asp2 transmembrane domain and the Asp2 protein, and where the polypeptide
 CC and the fragment retain the beta-secretase activity of the mammalian Asp2
 CC protein. Also included is an isoform of amyloid protein precursor (APP)
 CC comprising the amino acid sequence of a APP or its fragment containing an
 CC APP cleavage site recognisable by a mammalian beta-secretase, and further
 CC comprising two lysine residues at the carboxyl terminus of the amino acid
 CC sequence of the mammalian APP or APP fragment. The polypeptides are used
 CC for assaying for modulators of beta-secretase activity; identifying
 CC agents that inhibit the APP processing activity of human Asp2 aspartyl
 CC protease (Hu-Asp2); identifying agents that modulate the activity of Asp2
 CC ; and for reducing cellular production of amyloid beta (Abeta) from APP.
 CC Agents identified by the above methods are useful for treating
 CC Alzheimer's disease; and for identifying modulators of amyloid-beta
 CC (Abeta) peptide production, for use in designing therapeutics for the
 CC treatment or prevention of Alzheimer's disease. Probes and primers
 CC derived from Asp nucleic acid sequences are useful for detecting Hu-Asp
 CC nucleic acids in in vitro assays and in Northern and Southern blots. The
 CC present sequence represents the amino acid sequence of T7-caspase-caspase
 CC 8-human-Asp-2a delta TM construct which has a T7 tag, a caspase 8 leader
 CC sequence and cleavage site, and lacks the transmembrane domain. This
 CC construct was used for bacterial expression and purification of human
 CC Asp2a. (Updated on 11-SEP-2003 to standardise OS field)
 XX
 SQ Sequence 425 AA;
 Query Match 100.0%; Score 144; DB 4; Length 425;
 Best Local Similarity 100.0%; Pred. No. 1.8e-13;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GYVEMTVGSPPTQTLNLTVDGSSNFAV 28
 DB 46 GYVEMTVGSPPTQTLNLTVDGSSNFAV 73

Search completed: July 26, 2005, 16:44:20
 Job time : 166 secs

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